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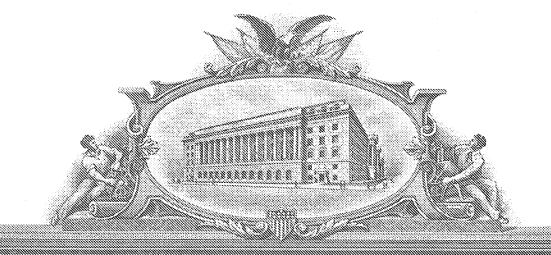
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UNITED STATES DEPARTMENT OF COMMERCE

United States Patent and Trademark Office

April 19, 2005

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PROVISIONAL APPLICATION FOR PATENT COVER SHEET

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c).

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City	Boston		State	MA	Zip	02110		
Country	US		Telephone	(617) 345-60	57 Fax	(617) 345-1	300	7
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This collection of information is required by 37 CFR 1.51. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 8 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden should be sent to the Chief Information Officer, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop Provisional Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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 $|\mathbf{X}|$ Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT

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Application Number	To be assigned	
Filing Date	Herewith	
First Named Inventor	C. Adra	
Examiner Name		
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Attorney Docket No.	732892-055040-P	

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1002 340	2002 170	Design filing fee		1401	330	2401	165	Notice of Appeal	
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1004 770	2004 385	Reissue filing fee		1403	290	2403	145	Request for oral hearing	
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: C

Chaker Adra

Application No.:

To be assigned

Group No.:

To be assigned

Filed:

Herewith Examiner:

To be assigned

For:

IDENTIFICATION OF GRANULOCYTE SUBTYPE-SELECTIVE RECEPTORS AND ION CHANNELS BY USING A HIGH-DENSITY

OLIGONUCLEOTIDE PROBE ARRAY

MAIL STOP PROVISIONAL APPLICATION Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

EXPRESS MAIL CERTIFICATE

"Express Mail" Label Number: EU 813933419 US

Date of Deposit: March 3, 2004

I hereby state that the following attached papers and fees:

- 1. Provisional Application for Patent Cover Sheet (1 pp.);
- 2. Provisional Patent Application (1423 pp.): Specification 18 pp.; Drawings 35 sheets; Attachment 1370 pp.;
- 3. Fee Transmittal (1 pg.);
- 4. Check in the Amount of \$80.00;
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are being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 C.F.R. section 1.10, on the date indicated above and is addressed to Box Provisional Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

Nicole M. Gignac

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IDENTIFICATION OF GRANULOCYTE SUBTYPE-SELECTIVE RECEPTORS AND ION CHANNELS BY USING A HIGH-DENSITY OLIGONUCLEOTIDE PROBE ARRAY

[0001] This invention was made in part with U.S. Government support under NIH Grant AI 43663 from the National Institute of Allergy and Infectious Diseases and by Grant RSG-01-241-01-LIB from the American Cancer Society (to C.A.). The U.S. Government has certain rights in this application.

FIELD OF THE INVENTION

[0002] The present invention relates to the identification of granulocyte subtype-selective receptors and ion channels that can be use as targets for drug discovery.

BACKGROUND OF THE INVENTION

[0003] Three types of human blood granulocytes, eosinophils, basophils and neutrophils, play roles in protecting against microbial infection by releasing cell type-specific mediators and proteases. Specifically, eosinophils and basophils evoke allergic reactions as well as damage nematodes. ^{1,2} As well as killing bacteria, neutrophils sometimes induce systemic vasculitis or multiple organ damage under certain conditions. ^{3,4} Thus, targeting granulocyte type-selective functions is considered an important strategy for drug discovery.

[0004] Activation of blood granulocytes and tissue mast cells is generally characterized by an influx of extracellular calcium (Ca²⁺), which is essential for subsequent release of granule-derived mediators, newly generated lipid mediators and cytokines. ⁵ The mechanism by which granulocyte mediator secretion is sustained is therefore likely to include modulation of various types of ion channels. Flow of ions including K⁺ and Cl⁻ may play an important role during granulocyte responses because they regulate cell membrane potential and thus influence Ca²⁺ influx. ⁶ Treatment of mast cells and basophils with pertussis toxin inactivates the Gi-type of G-proteins and abolishes degranulation induced by non-immunological ligands such as thrombin and N-formylpeptide; however, it fails to inhibit the influx of Ca²⁺. ⁷ Thus, Ca²⁺-independent

stimulation of Gi is also involved in granulocyte degranulation. The thrombin (protease) activated receptors and formylpeptide receptors are classified as G protein-coupled receptors (GPR), having an extracellular N-terminal segment, a seven transmembrane region, which forms the transmembrane core; three exoloops, three cytoloops, and a C-terminal segment. ⁸ Thus, ion channels and GPR both play essential roles in degranulation as well as other cellular function important for granulocytes. As a result, both ion channels and GPR are targets of drug development. ⁹

[0005] As the human genome project nears completion, the identification of potential drug targets using gene expression profiles from specific cell types is becoming practical and important for drug discovery. ^{10,11} The sequencing of the human genome is offering an unprecedented opportunity for the pharmaceutical development of drugs. Receptor genes and ion channel genes are found only in 5% and 1.3% of all genes present in the human genome, ¹⁰ respectively. However, receptors and ion channels are respectively found in 45% and 5% of the molecular targets of all known drugs. ^{9,12,13} Thus, receptors including GPR and ion channels are now considered as the most important drug targets.

[0006] Until recently, it has been impractical to analyze genome-wide expression of leukocytes. Newly developed technology, the microarray or high density oligonucleotide probe array (GeneChip) is one of the latest breakthroughs in experimental molecular biology, which allows approximately 39,000 transcripts derived from a cells transcriptome to be simultaneously monitored. Using this technology, we previously reported the transcriptome profiling of various types of mast cells and eosinophils. ¹⁴⁻¹⁶ However, there is still a need in the art to identify drug targets that are selectively, or preferentially, expressed in specific cell types such that efforts required for pharmaceutical development are minimized.

SUMMARY OF THE INVENTION

[0007] In the present study, we used GeneChip (version U133A containing approximately 22,000 gene probes) to examine the cell type-selective transcriptome expression of 7 types of leukocytes (basophils, eosinophils, neutrophils, CD4⁺ cells, CD8⁺ cells, CD14⁺ cells and CD19⁺ cells), platelets, mast cells and fibroblasts by focusing on the expression of granulocyte-selective genes for ion channels, GPR and other receptors. We identified many novel granulocyte subtype-selective transcripts that are useful for drug development.

[0008] Granulocyte subtype selective transcripts were chosen based on several conditions such as the transcript having 5-fold or greater expression level compared to the maximum level of other leukocytes. Fifty-one transcripts were chosen to be preferentially expressed by each granulocyte subtype. Seventeen out of the 51 transcripts have not been previously reported as granulocyte subtype-selective. Among the 17 receptors and ion channels, six were basophiland/or eosinophil-selective and were not highly expressed by other organs, indicating that they may be potential targets for anti-allergic drugs.

[0009] Utilization of this database (Attachment A) of potential cell type-selective drug targets will minimize the efforts required for pharmaceutical development of drugs for treatment of diseases of the immune system, cancer, cardiac diseases, as well as other diseases.

BRIEF DESCRIPTION OF THE DRAWINGS

[00010] Figure 1 shows Real-time quantitative PCR analysis of granulocyteselective gene expression. The relative mRNA expression level by each cell type against PBMNC was shown after normalization of mRNA levels for A. HTm4 (0.42 per 100 GAPDH), B. Ca²⁺ receptor alpha 1D subunit (0.003), C. prostaglandin E receptor type 3a2 (0.12), D. EMR-1 (0.62), and E. aquaporin 9 (0.92) expressed by PBMNC. Ne; neutrophils (n=3), Eo; eosinophils (n=2), Ba, basophils (n=3), CD4; CD4+ cells (n=3), P; PBMNC (n=1).

[00011] Figure 2 shows the demonstration of HTm4 protein on human basophils. Cells on the glass slide were incubated with 2 μg/ml polyclonal rabbit anti-hHTm4 antibody or 2 μg/ml rabbit IgG (H+L) as a control followed by incubation with a secondary antibody, highly cross-adsorbed Alexa Fluor® 546 conjugated goat anti-rabbit IgG (H+L) (Red). After mounting using the Prolong AntiFade Kit, slides were scanned by Zeiss Laser Scanning Microscope 5 Pascal.

[00012] Figures 3A through 3E shows granulocyte subtype-specific transcripts for ion channels and receptors.

[00013] Figures 4A through 4F show a table of "normalized AD" expression levels of various genes in indicated cells and shows corresponding graphs; x axis represents "normalized AD" expression levels.

[00014] Figures 5A through 5D show a table of "normalized AD" levels of various genes in indicated cells and shows corresponding graphs; x axis represents "normalized AD" expression levels.

[00015] Figures 6A through Figure 6R show the complete list of granulocyte subtype-selective transcripts.

DETAILED DESCRIPTION OF THE INVENTION

[00016] In this study, we have used high density oligonucleotide probe array (GeneChip) to measure the expression levels of approximately 20,000 different transcripts in highly purified cells. These cells were basophils, eosinophils, neutrophils, monocytes (CD14⁺), T lymphocytes (CD4⁺ and CD8⁺ cells), B lymphocytes (CD19⁺), lung-derived mast cells, cord blood-derived cultured mast cells, and nasal polyp-derived fibroblasts. The GeneChip assay allows the simultaneous measurement of large numbers of transcripts using relatively small numbers of cells. Using this technology, we could even measure triplicate transcriptome levels of basophils, the most rare granulocytes in peripheral blood.

[00017] Cell type-selective transcripts were selected based on the following criteria; (1) the average "normalized AD" expression level of each gene in a certain cell type must be 3-fold or greater than the maximal level in other cell types, and (2) must be significantly (p < 0.01) greater than that in other cell types. (3) The "AD" expression level provided with "absence" or "marginal" call by GeneChip Suite Software should be observed only once or not at all in the three or four independent experiments (3 experiments for basophils, 4 experiments for eisinophils and neutrophils) using different cell populations performed. (4) For the transcripts preferentially expressed for the two different cell types such as basophils and eosinophils, the average "normalized AD" expression levels in the two cell types should be within 3-fold of each other. Using these standards, we found 83 basophil-selective, 37 eosinophil-selective, 257 neutrophil-selective, 34 basophil-eosinophil-selective, 19 eosinophil-neutrophil-selective, and 17 basophil-neutrophil-selective transcripts. Due to the functional similarity with basophils, mast cell-selective transcripts were also examined, and 63 mast cell-selective and 11 mast cellbasophil-selective transcripts were also detected (Table 1, and Figures 6A-6R). Since mast cells, basophils and eosinophils play similar roles in allergic inflammation, the transcripts preferentially expressed for the three granulocytes by comparing their average "normalized AD"

levels to other leukocytes. Thirty-four transcripts were then selected; however, most of them were overlapped with the transcripts listed in Figures 6A-6R. Only four transcripts (MYB, SAMSN1, BACE2, and CASP3) were found not overlapped, and they were not receptors or ion channels.

[00018] Among the 491 granulocyte-selective transcripts listed in Figures 6A-6R, 4 ion channels, 19 GPR and 28 other receptors were further selected (Figure 3). When plural transcripts obtained by different probe sets had identical Genebank or Unigene accession numbers (http://www.ncbi.nlm.nih.gov/), the transcript showing the highest expression level was selected.

Ion channels and receptors preferentially expressed by granulocytes other than neutrophils

[00019] Eosinophils, basophils and mast cells play an important role in the pathogenesis of allergic diseases, but do not play an essential role in killing microbes except for nematodes. On the other hand, neutrophils play a crucial role in killing microbes such as bacteria. Caution should be taken in regulating neutrophil function even in the case of neutrophil-induced inflammation. Therefore, the molecules present only in granulocytes except for neutrophils would be important pharmaceutical targets for allergic disorders. ^{1,2}

[00020] Among the 51 granulocyte-selective transcripts for ion channels and receptors, we identified 17 granulocyte-selective transcripts that have not been reported for their selective expression (shown in bold letters in Figure 3). Of these 17 transcripts, eight were preferentially expressed by granulocytes other than neutrophils. Among these eight transcripts, the two transcripts for fibroblast growth factor receptor 2 and low density lipoprotein receptor were found to be expressed by multiple tissue cell types (shown at the Web site http://www.lsbm.org/index_e.html), which displays genomic expression of 55 different human tissue cells such as brain, heart and lung cells using the same experimental system. Affymetrix, U133A as ourts. Thus, they may not be suitable as a drug target because important organs that are unrelated to allergic inflammation (such as the brain) express it. Among the six novel transcripts found to be preferentially expressed by granulocytes except for neutrophils, we focus on the following four transcripts expressed by granulocytes including basophils. They were Ca²⁺ channel (*CACNA1D*), a prostaglandin E receptor, (*EP3A2*), epidermal growth factor-like module-containing mucin-like receptor (EMR) 1 (*EMR1*), and HTm4 (*MS4A3*).

[00021] Basophils are the rarest granulocytes present in human peripheral blood and as such their complete transcriptional profiles remain unclear and no basophil-selective transcripts have previously been reported. ²¹ Although eosinophils and mast cells have been considered as important therapeutic targets for allergic diseases for a long time, recent studies suggest the importance of basophils in pathogenesis of severe allergic diseases such as fatal asthma. ^{2,22} Therefore, we further examined the selective expression of these four basophil-, or basophil-eosinophil-selective transcripts by using real-time RT-PCR. As shown in the Fig. 1, including a known molecule preferentially expressed by neutrophils, aquaporin 9, ²³ the results obtained with GeneChip assay were confirmed by using this highly accurate and reproducible method. ²⁴

[00022] Among these four molecules, we could obtain a suitable antibody against HTm4, which is a member of a family of four transmembrane- proteins which include CD 20 and high affinity Fc receptor for IgE (Fc ϵ RI) β -chain. ²⁵ Genetics provided evidence for the existence of multiple loci relevant to atopic asthma on chromosome 11q13, including HTm4. ²⁶ Most recently, we have published data identifying HTm4 as a hematopoietic cell cycle regulator. ²⁷ Using specific antibody against HTm4, we could detect the expression of HTm4 at the protein level in basophils (Fig. 2). To confirm whether these ion channels and receptors could be potential drug targets for diseases involving basophil activation, the amount of molecules expressed by various cell types should be quantified and the effect of any identified antagonists should be tested on the cell types found to express these molecules.

[00023] As has been well documented and expected, Fc ε RI α , IL-3 receptors, IL-5 receptors, chemokine receptor CCR3, ^{1,2} sialic acid binding Ig-like lectin (Siglec)-8, ²⁸ Siglec-6, ¹⁵ histamine H4 receptor ²⁹ and chemoattractant receptor- homologous molecule expressed on Th2 cells (CRTH2) ³⁰ were preferentially expressed by basophils and/or eosinophils or mast cells. These consistent observations strengthened the reliability of the present methods and research strategy. Indeed, the antibody against Siglec-8 can induce selective apoptosis of eosinophils, and is expected to be useful therapeutically. ²⁸

Ion channels and receptors preferentially expressed by granulocytes including neutrophils

[00024] Of the 17 transcripts that have not been reported as granulocyte-selective, nine were preferentially expressed by granulocytes including neutrophils. Pharmaceutical targets

of selective granulocyte transcripts should treat inflammatory diseases without affecting the function of important organs that are unrelated to inflammation as well as the function of lymphocytes. However, four of the nine neutrophil-selective transcripts were expressed by multiple organ tissues. One of the four neutrophil-selective transcripts encoded proteinase-activated receptor (PAR)-2, a receptor for mast cell tryptase, which is linked to the pathogenesis of allergic diseases such as asthma. ^{31,32} PAR-2 transcripts are also abundantly expressed by tissue types including skin and intestine which are often the target organs for allergic diseases. But the development of PAR-2 antagonists for use as anti-allergic drugs may be unsuitable because it may down regulate neutrophil function and thereby induce bacterial infection.

General discussion

[00025] We identified 51 granulocyte-selective genes for ion channels and receptors by examining approximately 20,000 kinds of transcripts derived from 16,000 genes from 10 different types of cells using U133A GeneChip, which covers approximately half of the genes present in the human genome. The majority of these transcripts encoded molecules known or expected to be granulocyte subtype-selective such as the IL-3 receptor and Fc ε receptors.

[00026] Mast cells expressed low levels of Fc ϵ RI α compared to basophils, and that even neutrophils expressed a substantial level of the receptor (Figure 3). This raises the possibility that GeneChip assay may not be suitable for detecting selective molecules. In the present study, however, only the GeneChip data obtained using cord blood-derived mast cells and lung mast cells could be employed due to the strict data selection based on the RNA quality (see Methods). As has bee reported, ^{14, 33} peripheral blood-derived cultured mast cells or skinderived mast cells expressed approximately 10-fold Fc ϵ RI α mRNA compared to cord blood-derived mast cells (data not shown). Also, as shown in Figures 6A-6R, only 2 of the 4 neutrophil samples expressed Fc ϵ RI α mRNA. This may be explained by the observation that only neutrophils obtained from some allergic donors express the molecule. ³⁴

[00027] We unexpectedly found 17 granulocyte-selective transcripts including HTm4. Basophil- and/or eosinophil-selective transcripts identified in our study could be potential therapeutic targets for allergic diseases because these granulocytes play a crucial role in allergic inflammation. ^{1,2} Granulocyte-selective transcripts could also be drug targets for other inflammatory diseases such as systemic vasculitis. ^{3,4} Analysis of cell type-selective transcripts

from database searches is expected to minimize the efforts required for drug discovery. The public database (http://www.lsbm.org/index_e.html) shows that some granulocyte-selective transcripts (18 out of 51) detected in our study are abundantly expressed by multiple (more than 3) organ tissue cell types using the same GeneChip U133A probe array. Thus, the safety of any candidate drug must be evaluated by comparing its efficacy (on granulocytes) with its toxicity (to organs). Six out of the 17 novel granulocyte-selective molecules may be excluded from drug development due to their expression in multiple organs unrelated to the diseases. Thus, our approach has identified 11 receptors and ion channels with therapeutic potential. Especially, among the 11 receptors and ion channels, seven were basophil- and/or eosinophil-selective and were not expressed by other organs, indicating that they may be potential targets for anti-allergic drugs.

[00028] Finally, it should be stressed that basophils, the rarest leukocytes, have recently been found to play a more crucial role than we ever proposed in the pathogenesis of intractable allergic diseases such as fatal asthma.^{35,36} Thus, targeting basophil receptors and ion channels such as HTm4 and Ca²⁺ channel CACNA1D is particularly expected for the future drug discovery. The importance of molecules known to be expressed by basophils may be reevaluated regarding its selectivity. Freshly-isolated resting basophils expressed the highest level of IL-4 compared to other cell types. Because the basophil purification procedure requires more isolation steps, ex vivo manipulation may activate the cells. However, it should also be noted that basophils have been recently found as the major source of IL-4 at least in asthma models.^{37,38}

Materials and Methods

Purification of leukocytes

[00029] All human subjects in this study provided written, informed consent, and the Ethical Review Boards at the relevant hospitals (National Center for Child Health and Development, and Jikei University School of Medicine) approved the study. The subjects used in this study were all healthy volunteers, especially having no allergic diseases.

[00030] Granulocytes and mononuclear cells were separated from venous blood of normal volunteers. Human basophils were semipurified by means of Percoll (Pharmacia, Uppsala, Sweden) density gradient centrifugation, and the cells were further purified by negative selection through use of a MACS Basophil Isolation Kit (Miltenyi BioTech, Bergisch-Gladbach,

Germany), as described previously. ¹⁷ Eosinophils were isolated by using Percoll (1.090 g/mL) density centrifugation. The eosinophils were further purified by negative selection with anti-CD16-bound micromagnetic beads, as described previously. ¹⁸ Neutrophils were isolated by using Percoll (1.085 g/mL) density centrifugation and further purified by negative selection using anti-CD81 antibody and antimouse IgG-bound micromagnetic beads to eliminate contaminating eosinophils. These granulocytes purified from human peripheral blood were spun down onto slide glass by Cytospin II (Shandon Southern Instruments, Inc., Sewickley, PA). The purity of these cells was evaluated based on 500 cells stained with May-Grünwald and Giemsa solutions.

[00031] For preparation of lymphocytes and monocytes, peripheral blood mononuclear cells (PBMNC) were isolated by centrifugation on lymphocyte separation medium (Organon Teknika Corp., Durham, NC). Monocytes (CD14⁺ cells) were prepared using magnetic beads-conjugated CD14⁺ antibody (CD14 MicroBeads; Miltenyi Biotec) from PBMNC. CD4⁺ and CD8⁺ cells were also respectively sorted using magnetic beads-conjugated CD4⁺ (CD4 MicroBeads; Miltenyi Biotec) and CD8⁺ antibodies (CD8 MicroBeads; Miltenyi Biotec) from PBMNC after depletion of CD14⁺ cells with MACS CD14 MicroBeads (Miltenyi Biotec). The purity of CD4⁺, CD8⁺ and CD14⁺ cells was evaluated by staining the magnetic beads- conjugated cells compared to feasible control cell preparations such as unpurified cells with FITC-labeled goat anti-mouse Immunoglobulin (BD Pharmingen, Tokyo, Japan). Peripheral B cells were purified by a combination of negative (MicroBeads-conjugated antibodies to CD3, CD7, CD14, CD42b, and CD56; Miltenyi Biotec) and positive (CD19 MicroBeads; Miltenyi Biotec) selection using MicroBeads (Miltenyi Biotec). To obtain platelet rich plasma, blood samples were mixed with 3.8% (w/v) sodium citrate solution (9:1) and centrifuged at 260x g for 15 min. at 20°C. To remove any contaminating erythrocytes and leukocytes, the plasma was centrifuged again at 260x g for 15 min.

[00032] Human mast cells were derived from cord blood CD34⁺ progenitor cells as described previously. ¹¹⁻¹⁴ Briefly, progenitor cells purified from peripheral blood by CD34⁺ isolation kits (Miltenyi Biotec), were cultured in Iscove's modified Dulbecco medium supplemented with 1% insulin-transferrin-selenium supplements (Life Technologies), 50 µM 2-mercaptoethanol, antibiotics, and 2% fetal calf serum in the presence of 100 ng/ml stem cell

factor and 50 ng/ml IL-6. After 11 to 14 weeks of culture, tryptase positive cells represented more than 99% of the cells.

Purification of human lung mast cells and nasal polyp-derived fibroblasts

[00033] Normal human lung tissue dissected during surgery was obtained macroscopically after informed consent. Human lung mast cells were dispersed from chopped lung specimens by an enzymatic procedure and were purified by magnetic bead affinity selection using the mAb anti-kit, YB5.B8 (BD PharMingen, San Diego, CA) as described previously. ¹⁹ The cells were further cultured in the presence of SCF and interleukin 6 (IL-6) for several weeks. Human nasal polyp-derived fibroblasts were obtained as previously reported. ²⁰

GeneChip expression analysis

Human genome-wide gene expression was examined using the Human [00034] Genome U133A probe array (GeneChip, Affymetrix, Santa Clara, CA), which contains the oligonucleotide probe set for 22,000 full-length genes. Experiments were performed in accordance with the manufacturer's protocol (Expression Analysis Technical Manual) and previous reports. 11-14 Total RNA (3-10 µg) was extracted from 10⁷ cells. Double-stranded cDNA was synthesized using a SuperScript Choice system (Life Technologies) and a T7-(dT)24 primer (Amersham Pharmacia Biotech, Buckinghamshire, UK). The cDNA was subjected to in vitro transcription in the presence of biotinylated nucleoside triphosphates using a BioArray high-yield RNA transcript labeling kit (Enzo Diagnostics, Farmingdale, NY). The biotinylated cRNA was hybridized with a probe array for 16 h at 45°C. In some experiments as indicated in the supplementary table, biotinylated cRNA was prepared using two-cycles of cDNA synthesis and in vitro transcription for target amplification was performed according to the manufacturer's "The Small Sample Labeling Protocol version II" (Affymetrix, Inc). For the latter protocol, we employed 100 ng total RNA. After washing, the hybridized, biotinylated cRNA was stained with streptavidin-phycoerythrin (Molecular Probes, Eugene, OR) and then scanned with a HP gene array scanner. The fluorescence intensity of each probe was quantified using a computer program, GeneChip Analysis Suite 5.0 (Affymetrix). The expression level of single mRNA was determined as the average fluorescence intensity among the intensities obtained by 11 paired (perfect- matched and single nucleotide-mismatched) probes. If the intensities of mismatched

probes were very high, gene expression was judged to be absent, even if a high average fluorescence was obtained with the GeneChip Analysis Suite 5.0 program. The level of gene expression was determined as the average difference (AD) using the GeneChip software. Each AD level was then normalized by dividing it with the median value of 22,283 AD levels obtained in an experiment ("normalized AD" level).

Real-time reverse transcriptase (RT)-PCR

[00035] Total RNA was isolated using Isogen (Nippon gene, Tokyo, Japan) according to the manufacturer's instructions and quantified by measuring the absorbance at 260 nm. RNA was subsequently treated with DNase I (Life Technologies) reverse transcribed using Superscript II reverse transcriptase (Life Technologies). Real-time RT-PCR was performed 10 ng cDNA in 25 µl of final volume using the primers and probes supplied by "Assays-on-Demand Gene Expression system" (PE Applied Biosystem) according to the manufacturer's instructions. Measurement of gene expression was performed using the ABI PRISM 7700 Sequence Detector. The expression level of each gene was normalized to a GAPDH.

Staining of basophils with anti-HTm4

[00036] Basophils purified from human peripheral blood with Basophil Isolation Kit (Miltenyi Biotec) were spun down onto slide glass by Cytospin II (Shandon Southern Instruments Inc., Sewickley, PA). Cells were fixed with aceton for 1 minute and then blocked in goat serum in 50 mm TRS-Cl, pH 7.4 for 1 hour. Cells were further incubated for 2 hours with 2 μg/ml of the polyclonal antibody rabbit anti-hHTm4. Cells were then washed three times with PBS and incubated with a secondary antibody, highly cross-adsorbed Alexa Fluor® 546-conjugated goat anti-rabbit IgG (H+L) (Red) (Molecular Probes, Eugene, OR) for one hour. After three PBS washes, air dried cells were further mounted using the Prolong Anti-Fade Kit (Molecular Probes, Eugene, OR). Slides were scanned by Zeiss Laser Scanning Microscope 5 Pascal (Carl Zeiss Microimaging Inc, Thornwood, NY).

Purity and viability of the leukocytes, and RNA quality

[00037] We used leukocyte samples in this study only if the purity of each cell type was at least 98%, but there are >0.5% contaminated cells in any of the samples. We could

not evaluate the purity of CD19⁺ cells and platelets due to lack of feasible controls or methods. However, specific transcript markers for non-granulocytes (CD4, CD8, CD14, CD19, IgG, etc.) as well as granulocyte subtype-specific transcripts were reasonably expressed by each leukocyte type as shown in Figures 6A-6R. Regarding the viability, we qualified the RNA before GeneChip assay using Array Quality Metrics Comparisons Software (Affymetrix) as well as trypan blue staining (they were always >95% viable), since RNAse-rich granules derived from degenerating cells rapidly destroy RNA transcripts. Briefly, to evaluate the quality of RNA, the ratio of 3'-probe set and 5'-probe set of housekeeping genes were compared as shown in Figures 6A-6R. According to the above software's guidance, the ratio of >2:1 at standard sample (5µg total RNA) protocol and that of >10:1 at small sample (50 ng total RNA) protocol were recommended. As shown in Figures 6A-6R, the cells used in the present study had the appropriate ratios of 3'-probe set and 5'-probe set of housekeeping genes, suggesting that these cells were highly viable.

Statistical analysis

[00038] Since logarithmic "normalized AD" levels were normally distributed within each group, unpaired parametric Student's two-tailed *t*-test was employed to analyze the data on a logarithmic scale.

Table 1. Representative cell type-selective transcripts in granulocytes

Accession # a Transcript	S.I. b	Normalized AD level .
Basophil-selective		
NM 000589.1 IL-4	73.3	13.3
L35848.1 HTm4	38.2	132.1
BC005912.1 Fc ε RI α	12.7	218.9
Eosinophil-selective		
NM 001140.1 Arachidonate		
15-lipoxygenase	74.1	18.3
NM_024703.1 FLJ22593	19.1	29.1
NM_014442.1 Siglec-8	9.8	16.9
Neutrophil-selective		
NM_004633.1 IL-1 R,		
type II	127.9	51.5
U73191.1		
inward rectifier K ⁺ channel Kir1.3	107.5	98
NM_001557.1 CXCR2 (IL-8 receptor β)	39.3	105.2
Mast cell-selective		
AF206667.1 tryptase β	84.3	159.4
NM_001911.1 cathepsin G	51.5	72.1
BC005929.1 major basic protein	31.6	72.5
Basophil-eosinophil-selective		
M75914.1 IL-5R α	42.8	19.4(B), 29.3(E)
NM_004778.1 CRTH2	16.6	23.9(B), 38.1(E)
NM_001828.3 Charcot-Leyden		,
crystal protein	15.2	229.2(B), 198.6(E)
Eosinophil-neutrophil-selective		
NM_005306.1 GPR 43 (PAR1-like)	21.7	11.7(E), 32.9(N)
NM_004668.1 DHHC		
domain containing 18	6.6	16.2(E), 44.5(N)
Basophil-neutrophil-selective		
NM_016006.1 CGI-58 protein	5.8	12.6(B), 21.2(N)
Basophil-mast cell-selective		
NM_001870.1 carboxypeptidase A3	59.2	111.7(M), 137.3(B)
NM_002529.2 TRK neurotrophin receptor	34.7	3.1(M), 7 (B)
NM_000139.1 Fc ε RI β	21.2	22.2(M), 43.8(B)

a. The GenBank accession number (http://www.ncbi.nlm.nih.gov).

b. Selectivity index (S.I.) was calculated by comparing the "normalized AD" level in a cell type or of two cell types with the maximal gene expression level of the other 8 or 9 cell types. The complete list of the genes having >3 S.I. is shown in Supplementary Table 1.

[00039] Figures 6A-6R show the complete list of granulocyte subtype-selective transcripts. Selectivity index (S.I.) was calculated by comparing the "normalized AD" level of a cell type or of two cell types with the maximal gene expression level of the other 8 or 9 cell types including platelets (Pl), CD4⁺ cells (CD4), CD8⁺ cells (CD8), CD14⁺ cells (CD14), CD19+ cells (CD19) and nasal polyp-derived cultured fibroblasts (Fb). Transcripts having S.I. >3-fold were shown in Figures 6A-6R A-H. A. Basophil (Ba)-selective transcripts. B. Eosinophil (Eo)selective transcripts. C. Neutrophil (Ne)-selective transcripts. D. Mast cell (MC)-selective transcripts. E. Basophil and eosinophil-selective transcripts. F. Eosinophil and neutrophilselective transcripts. G. Basophil and neutrophil-selective transcripts. H. Mast cell and basophilselective transcripts. I. Raw AD levels for the median values used to normalize the raw AD levels, and the housekeeping genes. When the result was accompanied by presence call, it was shown as a bold numeral. Italic numerals show that the raw AD levels were associated with absence call by the GeneChip analysis software. 1. Abbreviations used in the tables were (small); the results obtained by the small sample protocol (see materials and methods), R; receptor, and ICN: ion channel.

REFERENCES

[00040] The references cited herein are incorporated by reference.

- 1. Bochner BS. Systemic activation of basophils and eosinophils: markers and consequences. J Allergy Clin Immunol. 2000;106:S292-302.
- 2. Hirai K, Miyamasu M, Takaishi T, Morita Y. Regulation of the function of eosinophils and basophils. Crit Rev Immunol. 1997;17:325-52.
- 3. Savage CO. The evolving pathogenesis of systemic vasculitis. Clin Med. 2002;2:458-64.
- 4. Oehler R, Weingartmann G, Manhart N, et al. Polytrauma induces increased expression of pyruvate kinase in neutrophils. Blood. 2000;95:1086-92.

- 5. Church MK, Pao GJ, Holgate ST. Characterization of histamine secretion from mechanically dispersed human lung mast cells: effects of anti-IgE, calcium ionophore A23187, compound 48/80, and basic polypeptides. J Immunol. 1982;129:2116-21.
- 6. Lin CS, Boltz RC, Blake JT, et al. Voltage-gated potassium channels regulate calcium-dependent pathways involved in human T lymphocyte activation. J Exp Med. 1993;177:637-45.
- 7. Saito H, Okajima F, Molski TF, Sha'afi RI, Ui M, Ishizaka T. Effects of ADP-ribosylation of GTP-binding protein by pertussis toxin on immunoglobulin E-dependent and -independent histamine release from mast cells and basophils. J Immunol. 1987;138:3927-34.
- 8. Ji TH, Grossmann M, Ji I. G protein-coupled receptors. I. Diversity of receptor-ligand interactions. J Biol Chem. 1998;273:17299-302.
- 9. Zambrowicz BP, Sands AT. Knockouts model the 100 best-selling drugs. Will they model the next 100? Nat Rev Drug Discov. 2003;2:38-51.
- 10. Venter JC, Adams MD, Myers EW et al. The sequence of the human genome. Science 2001;291:1304–51.
- 11. International Human Genome Sequencing Consortium. Initial sequencing and analysis of the human genome. Nature 2001;409: 860–921.
- 12. Drews J. Drug discovery: a historical perspective. Science. 2000;287:1960–4.
- 13. Hopkins AL, Groom CR. The druggable genome. Nature Rev Drug Discov. 2002;1:727–30.
- 14. Iida M, Matsumoto K, Tomita H, et al. Selective down-regulation of high-affinity IgE receptor (FcεRI) α-chain messenger RNA among transcriptome in cord blood-derived versus adult peripheral blood-derived cultured human mast cells. Blood. 2001;97:1016-22.

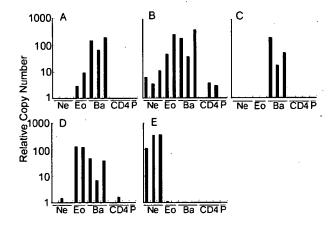
- 15. Nakajima T, Matsumoto K, Suto H, et al. Gene expression screening of human mast cells and eosinophils using high-density oligonucleotide probe arrays: abundant expression of major basic protein in mast cells. Blood. 2001;98:1127-34.
- Nakajima T, Inagaki N, Tanaka H, et al. Marked increase in CC chemokine gene expression in both human and mouse mast cell transcriptomes following Fcε receptor I cross-linking: an interspecies comparison. Blood. 2002;100:3861-8.
- 17. Yoshimura C, Miyamasu M, Nagase H, et al. Glucocorticoids induce basophil apoptosis.

 J Allergy Clin Immunol. 2001;108:215-20.
- 18. Matsumoto K, Schleimer RP, Saito H, Iikura Y, Bochner BS. Induction of apoptosis in human eosinophils by anti-Fas antibody treatment in vitro. Blood. 1995;86:1437-43.
- 19. Okayama Y, Hunt TC, Kassel O, Ashman LK, Church MK. Assessment of the anti-c-kit monoclonal antibody YB5.B8 in affinity magnetic enrichment of human lung mast cells. J Immunol Methods. 1994;169:153-61.
- 20. Fukagawa K, Okada N, Fujishima H, et al. CC-chemokine receptor 3: a possible target in treatment of allergy-related corneal ulcer. Invest Ophthalmol Vis Sci. 2002;43:58-62.
- 21. Arock M, Schneider E, Boissan M, Tricottet V, Dy M. Differentiation of human basophils: an overview of recent advances and pending questions. J Leukoc Biol. 2002;71:557-64.
- 22. Kepley CL, McFeeley PJ, Oliver JM, Lipscomb MF. Immunohistochemical detection of human basophils in postmortem cases of fatal asthma. Am J Respir Crit Care Med. 2001;164:1053-8.
- 23. Loitto V-M, Forslund T, Sundqvist T, Magnusson K-E, Gustafsson M. Neutrophil leukocyte motility requires directed water influx. J Leukoc Biol. 2002;71:212-22.
- 24. Pfaffl MW. A new mathematical model for relative quantification in real-time RT-PCR. Nucleic Acids Res. 2001;29:e45.

- 25. Adra CN, Lelias JM, Kobayashi H, et al. Cloning of the cDNA for a hematopoietic cell-specific protein related to CD20 and the beta subunit of the high-affinity IgE receptor: evidence for a family of proteins with four membrane-spanning regions. Proc Natl Acad Sci U S A. 1994;91:10178-82.
- 26. Adra CN, Mao XQ, Kawada H, et al. Chromosome 11q13 and atopic asthma. Clin Genet. 1999;55:431-7.
- 27. Donato JL, Ko J, Kutok JL, et al. Human HTm4 is a hematopoietic cell cycle regulator. J Clin Invest. 2002;109:51-8.
- 28. Nutku E, Aizawa H, Hudson SA, Bochner BS. Ligation of Siglec-8: a selective mechanism for induction of human eosinophil apoptosis. Blood. 2003;101:5014-20.
- 29. O'Reilly M, Alpert R, Jenkinson S, et al. Identification of a histamine H4 receptor on human eosinophils--role in eosinophil chemotaxis. J Recept Signal Transduct Res. 2002;22:431-48.
- 30. Hirai H, Tanaka K, Yoshie O, et al. Prostaglandin D2 selectively induces chemotaxis in T helper type 2 cells, eosinophils, and basophils via seven-transmembrane receptor CRTH2. J Exp Med. 2001;193:255-61.
- 31. Moffatt JD, Jeffrey KL, Cocks TM. Protease-activated receptor-2 activating peptide SLIGRL inhibits bacterial lipopolysaccharide-induced recruitment of polymorphonuclear leukocytes into the airways of mice. Am J Respir Cell Mol Biol. 2002;26:680-4.
- 32. Schmidlin F, Amadesi S, Vidil R, et al. Expression and function of proteinase-activated receptor 2 in human bronchial smooth muscle. Am J Respir Crit Care Med. 2001;164:1276-81.
- 33. Bradding P., Okayama Y., Kambe N., Saito H., Ion channel gene expression in human lung, skin and cord blood-derived mast cells, J. Leukoc Biol 2003; 73:614-20.

- 34. Gounni AS, Lamkhioued B., Koussih L., Ra C., Renzi PM, Hamid Q., Human neutrophils express the high-affinity receptor for immunoglobulin E (Fc epsilon RI): role in asthma. FASEB J. 2001; 15:940-9.
- 35. Koshino T., Teshima S., Fukushima N., Takaishi T., Hirai K., Miyamoto Y., et al. Identification of basophils by immunohistochemistry in the airways of post-mortem cases of fatal asthma. Clin Exp Allergy 1993; 23:919-25.
- 36. Kepley C.L., McFeeley P.J., Oliver J.M., Lipscomb M.F., Immunohistochemical detection of human basophils in postmortem cases of fatal asthma. Am J Respir Crit Care Med 2001; 164:1053-8.
- 37. Schroeder J.T., Lichtenstein L.M., Roche E.M., Xiao H., Liu M.C., Il-4 production by human basophils fund in the lung following segmental allergen challenge. J. Allergy Clin Immunol 2001; 107:265.71.
- 38. Luccioli S., Brody D.T., Hasan S., Keane-Myers A., Prussin C., Metcalfe DD, IgE(+), Kit (-), I-A/I-E-(-) myeloid cells are the initial source of Il-4 after antigen challenge in a mouse model of allergic pulmonary inflammation. J Allergy Clin Immunol 2001; 110:117-24.

Figure 1



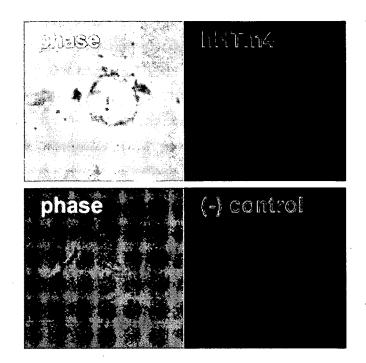


Figure 2

Figure 3. Granulocyte subtype-specific transcripts for ion channels and receptors

Ion Channels													
Transcript (Accession #, GenBank)	Cell-	MC	Ва	Eo	ž	귭	CD4	CD8	C D14	CO c	Fb	$\mathcal{L}_{\mathcal{C}}$	Gene Functions
Ca2+ channel type A1 D (BE550599)	rype Ba, Eo	0.1	1.7	1.5	0.4	0.0	0.3	0.1	0.0	0.3	0.0	0	
aquaporin 9 (NM 020980) 602914	Se Se	0.7	0.1	0.4	137.4	0.4	6.0	0.1	9.8	0.1	0.2		facilitates uptake of the metalloids arsenite and
													antimonite
K*channel Kir1.3 (U73191) 600359	Se	6.0	0.2	0.4	99.5	0.7	0.2	0.0	8.0	0.0	0.0	S	Andersen syndrome (170390)
K ⁺ channel Kir2.1 (AF153820) 600681	Ž	0.7	80	5.3	40.7	0.4	0.2	0.5	. 5.1	0.7	-	9	and Bartter syndrome (241200) Andersen syndrome (170390)
	<u>!</u>	;	2	:			ļ <i>.</i>	<u>.</u>					and Bartter syndrome (241200)
GPCR histamine H ₄ R (<u>AF312230) 606792</u>	Ba	0.7	34.2	9.4	0.7	0.4	8.0	0.5	. 9.0	0.0	0.1	0	expression of HRH4 conferred
TOOTE COTOCOX PROFILE TO THE STANDARD OF THE S	å	ò		-	,	70	ć	ç		9		•	sensitivity
73.8 (1162027) 605246	Ba Ba Fo	0.0	55.7	39.4	7.0	0.0	0.5 1	7.0	3.0	0.0	. 0	o ~	signalling paulways ananhylatoxin recentor
CCR3 (NM_001837) 601268	Ba, Eo	9.0	117.	6.06	24.9	0.2	0.5	0.4	0.2	0.2	0.4	0	importance for eosinophil
			4	. ;	,	,			,	,		,	responses
CRTH2 (NM 004778) 604837	Ba, Eo	Ξ:	26.0	38.2	2.0	8.0	4.	0.1	1.2	6.0	0.5	0	the cell via activation of
						•							rotrimeric G proteins
EMR-1 (NM 001974) 600493	Ba, Eo	8.0	33.5	6.06	·4.2	3.4	1.7	8.0	7.1	1.6	0.5	1	Probably involved in cellular
	I	,		;			,			,	(,	response to a hormone
adenosine A, R (NM 000677) 600445	8 9	2.6	2.9	15.4	2.3	1.2	9.1	0.5	2.0	0.5	0.5	o·0	cardioprotective function P2BV2 may participate in
600041	3	-		Ç	-	7:0	1.0		7: -	7.0	- -	2	of the cell cycle
													endometrial carcinoma cells
GPRIOSparinergick (NM 014879)	E E	5.6	2.9	15.4	2.3	1.2	9.1	0.5	2.0	0.5	0.5	7	GPR105 is a G-protein-coupled
													dentifying a quiesce
												:	primitive population of
													pone marrow; Orkitos impilia
													heral and neuroimmu
GPR, Edg-4 (AF011466) 605110	Eo, Ne	1.3	2.8	15.9	24.2	0.1	3.6	5.0	3.8	1.9	6.0	7	edg-4 mma was expressed in
													mouse islets; edg-4 (lpa2) r is a
													티
													ovarian carcinoma, and is
			٠										essed po
													extended gain-of-function
PAR1-like GPR43 (NM 005306)	Eo, Ne	0.3	0.7	12.4	35.2	8.0	0.1	0.1	0.7	0.4	0.1	. 0	the highest levels of gpr43 were
		٠.						٠					

found in immune cells; gpr43 is highly restricted in hamatonoisit itsense	receptor for the chemotactic and inflammatory peptide anaphylatoxin c5a. this receptor stimulates chemotaxis, granule enzyme release and superoxide anion production.	receptor to interleukin-8, which is a powerful neutrophils chemotactic factor. binding of il-8 to the receptor causes activation of neutrophils. this response is mediated via a g-protein that activate a phosphatidylinositol-calcium second messenger system. this receptor binds to il-8 with a high affinity and to mgsa (gro) with a low	receptor to interleukin-8, which is a powerful neutrophils chemotactic factor. binding of il-8 to the receptor causes activation of neutrophils. this response is mediated via a g-protein that activate a phosphatidylinositol-calcium second messenger system; this receptor binds to il-8 with a high affinity and to gro/mgsa and nap-2 also with a high affinity with a high affinity	HIGH AFFINITY FOR METHIONYL METHIONYL NEUTROPHILS CHEMOTACTIC RECEPTOR RECEPTOR CAUSES ACTIVATION NEUTROPHILS RESPONSE IS MEDIATED
	~	0	· ~	0
	9.4	1.0	0.0	0.5
	1.0	0.1	9.0	2
	25.6		0.7	62.6
	0.5	0.2	8.0	0.7
	2 .	0.3	0.3	£.
	2.3	4.0	2.7	3.6
	92.6	4.	112.1	282.9
	13.6	0.3	.	∞ ∞
	21.6	4. E	Ė	23.5
	2.3	0.2	0.2	5.8
	Se .	ž	÷	ž
603823	CSa R (<u>NM_001736) 113995</u>	CXCR1 IL-8R (NM 000634) 146929	CXCR2 IL-8R (<u>NM_001557)</u>	formyl peptide R I <u>(NM_002029)</u> <u>136537</u>

VIA A G-PROTEIN THAT ACTIVATES PHOSPHATIDYLINOSITOL- CALCIUM SECOND MESSENGER SYSTEM	5 likely FPR2 mediates superoxide production at high concentrations of fMLF	2 CSL2 is an anaphylatoxin- binding protein with unique ligand binding and signaling	GPR86 proved to be a G(i)-coupled receptor displaying a high affinity for ADP, similar to the P2Y(12) receptor and can therefore be tentatively called P2Y(13)	15 PAR2 plays a key role in chronic joint inflammation	0 responsible for initiating the allergic response	I Cell cycle regulator	0 promotes the proliferation and differentiation of hematopoietic cells	 engagement of 2B4 with specific antibody activates NK cytolytic activity 	17 Receptor for acidic and basic fibroblast growth factors.	0 lacking either 115ra or Sox 4 have defects in B-cell development	0 SIGLEC8 expression on eosinophils but not other leukocytes	Signaling from the KIT receptor tyrosine kinase is essential for primordial germ cell growth	OBBP1 is almost exclusively expressed on B cells.	0 MS4A2 Allergic disease 34 receptor-mediated endocytosis
	0.1	0.3	0.0	1.2	0.2	0.0	0.1	0.1	1.0	0.1	0.2	0.5	0.0	6.3
	0.8	0.1	0.6	0.4	. 0.4	0.2	0.1	Ξ.	0.1	0.2	0.5	0.1	0.4	0.3
	0.9	1.0	12.3	1.6	<u>4</u> .	0.1	0.7	5.2	0.1	0.1	0.2	0.1	0.0	3.5
	9.0	0.2	.0.2	0.1	8.0	0.4	0.2	3.5	0.2	0.1	0.1	0.5	0.0	3.4
	1:0	0.3	0:1	0.7	5.1	0.5	0.3	0.4	0.1	0.1	0.2	0.2	0.0	0.5
	0.1	0.0	1.9	0.1	4.	0.2	0.2	0.1	0.1	0.1	0.3	0.8	0.5	1.6
	75.5	3.4	88.2	36.2	5.5	3.5	0.7	1.2	0.2	9.0	9.4	1.2	0.2	0.6
	9.0	0.8	17.9	2.2	4.6	6.9	5.0	16.5	12.1	30.4	17.4	4.2	0.0	0.4
•	0.5	6.0	0.2	0.3	219.	133.	52.6	56.0	27.9	20.5	0.3	7.2	0.2	44.3
	0.4	0.1	0.3	0.1	19.4	9.0	9.0	0.3	0.1	6.0	8:1	89.0	5.6	22.3
	Ne S	Š	ž	S S	Ba	Ba	Ba	Ва, Ео	.Ba, Eo	Ва, Ео	E0	MC	MC.	MC, Ba MC, Ba
	formyl peptide R 2 (<u>U81501</u>)	GPR77 (NM_018485)	GPR86 purinergic R (<u>NM 023914)</u>	PAR2 (BE965369) 600933	<u>Other Receptors</u> Fc.ε RI α (<u>BC005912) 147138</u>	HTm4 (L.35848) 606498	IL-3 R (<u>NM_002183) 308385</u>	GD2431NK/ceiiR (NM_016382) 605554	fibroblast growth factor R 2 (NM 022969) 176943	IL-5R α (M75914) 147851	Siglec 8 (<u>NM_014442) 605639</u>	CD117 c-KIT (<u>NM_000222) 164920</u>	Sigles 6 (D86.358) 604405	Fc & RIB (NM 000139) 147138 low density lipoprotein R

participates in the primary signal transduction mechanism of NGF; is also an immunoregulatory cytokine acting on monocytes	his gene is mainly expressed in small intestine, colon, testis, and leukocytes	Receptor for TNFSF2/TNF- alpha and homotrimeric TNFSF1/lymphotoxin-alpha. The adaptor molecule FADD recruits caspase-8 to the activated receptor. The resulting death- inducing signaling complex (DISC) performs caspase-8 proteolytic activation which initiates the subsequent cascade of caspases (aspartate- specific cysteine proteases) mediating apoptosis. Contributes to the induction of noncytocidal TNF effects including anti-viral state and activation of the acid submonedinase.	Receptor for TNFSF6/FASL. The adaptor molecule FADD recruits caspase-8 to The activated
<i>o</i>	1		7
0.0	1.3	9	9.7
0.0	5.1	9.0	2.6
0.1	6.0	17.4	3.6
0.1	1.6	6.6	8.
0.1	1.7	2.2	5.6
0.1	<u>5.</u>	<u>.</u>	1.3
0.0	6.9	74.7	50.4
0.1	1.5	7.8	7.3.
7.3	1.7	7.7	10.9
4.6	Ξ	1.7	<u>4</u> .
MC, Ba	e N	e Z	Š
(NM_000527) <u>606945</u> TRK neurotrophin R (NM_00252 <u>9)</u> 19131 <u>5</u>	butyrophilin like R (<u>AK025267)</u>	CD120a, TNF-R-I (<u>NM_001065</u>)	CD95, Fas, APO-1 (<u>AA164751) 134637</u>

activation which initiates the subsequent cascade of caspases (asparate-specific cysteine proteases) mediating apoptosis. FAS- mediated apoptosis may have a role in the induction of peripheral tolerance, in the antigen-stimulated suicide of mature T-cells, or both. The secreted

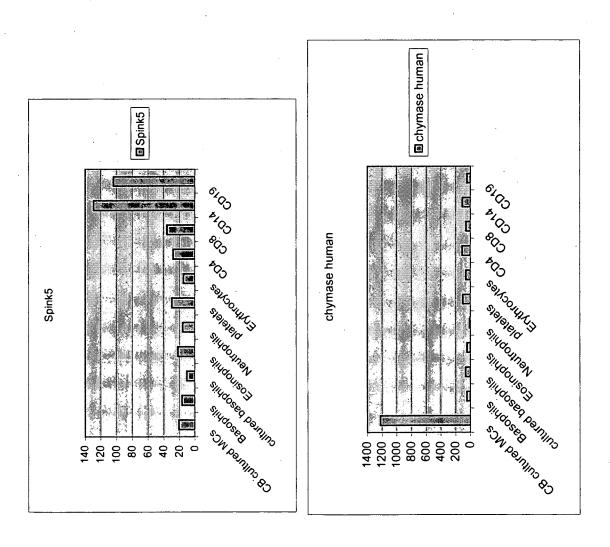
receptor. The resulting deathinducing signaling complex (DISC) performs caspase-8

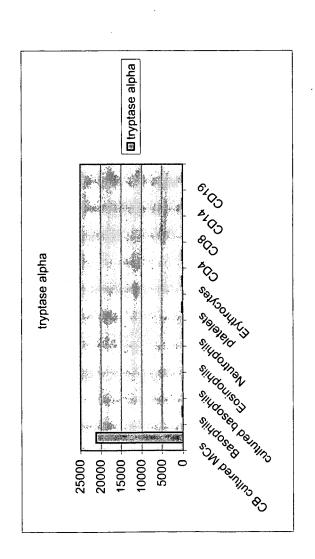
proteolytic

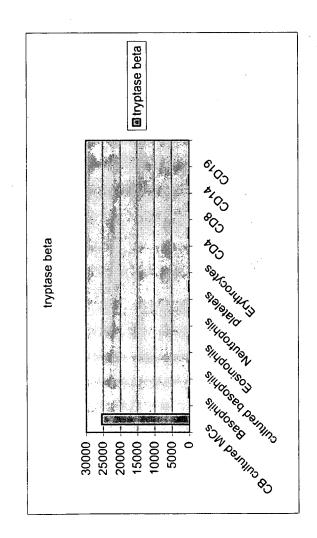
isoforms 2 to 6 block apoptosis (in vitro). does not induce apoptosis.		failure of the patient's neutrophils to express Fc receptor III was associated with SLE	inhibits cell proliferation and survival in response to CSF3	romotes the proliferation and differentiation of hematopoietic cells	type II receptor inhibits IL.1 activity by acting as a decoy target for IL.1	IGFs may elicit a myogenic event which may be a key mechanism in the etiology of breast and ovarian cancer	Mutation in either MPRD or MPRI might result in a clinical disorder resembling a mucolipidosis	Eosinophils may be activated through LIR7 for release of eosinophil-derived neurotoxin	that TLR2-TLR1 heterodimers mediated the strongest cell activation	TLR2 is a molecular link between microbial products, apoptosis, and host defense mechanisms	immunostimulatory activity
0	2 5	4	₩.	0	æ	~	32	7	0	0	0
. 9.0	0.2	0.1	0.2	T , .	0.0	3.5	15	0.1	0.3	0.4	9.0
0.1	2.3	2.6	. 0.1	1.3	0.0	2.0	8.4	0.5	5.	6.0	1.2
0.7	7.2	2.2	25.5	2.9	0.2	2.4	8.4	11.6	3.0	26.3	2.1
0.1	0.1	1.3	0.2	0.3	0.1	2.8	8.7	0.1	0.7	0.1	6.0
0.2	0.2	1.3	8.0	0.4	0.4	0.1	2.9	0.0	. 0.8	13	0.9
0.5	0.3	9.9	0.2 '	0.4	0.2	9:1	1.7	1.8	1.6	1.6	0.1
78.7	59.9 84.3	199.6	163.6	14.1	53.5	17.4	85.3	41.2	31.5	83.8	8.8
7.7	7.8	6.1	1.6	2.0	0.1	5.0	5.4	4.3	1.2	1.3	6.0
1.5	2.4	9.1	0.4	0.2	0.1	3.5	6.0	5.8	0.3	0.9	1.0
0.1	1.5	0.7	0.1	0.3	0.1	0.3	4.6	0.5	9.0	6.0	0.5
N.	S S	N G	Ne	Š S	Š	Še	Ne Se	Ne	Š	Ne	Se Se
decoy R1, TRAILR3 (<u>AF012536)</u>	<u>8038013</u> Fcy R IIc2 (<u>U90939)</u> Fcy R IIc3 (U90940)	Fcy R III (<u>104162) 146740</u>	G-CSF R (NM 0007601) 138971	IL-13 R (<u>U81379) 308385</u>	IL-1R, type II (<u>NM_004633) 147811</u>	IGFR I (<u>NM_000875) 147370</u>	IGFR 2 (<u>NM_000876) 147280</u>	leukocyte immunoglobulin-like R A2 (NM_006866) 604812	Toll-like R 1 (<u>AL050262) 601194</u>	Toll-like R 2 (<u>NM_003264) 603028</u>	Toll-like R 6 (NM_006068)

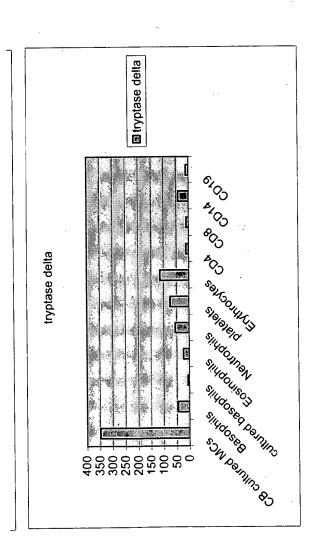
a. Cell-type specificity was obtained by comparing the "normalized AD" levels of each gene in mast cells (MC; average of 2 experiments), basephils (Ba; average of 3 experiments), average of 4 experiments), neutrophils (Ne; average of 4 experiments), platelets (PI), CD4* cells (CD4), CD8* cells (CD8), CD14* cells (CD19+ cells (CD19) and nasal polyp-derived cultured fibroblasts (Fb).

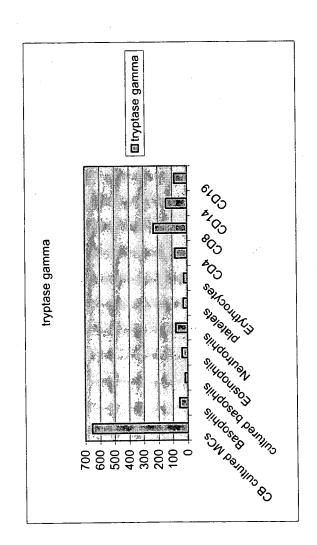
	CB cultured MCs Br	Roconhile F	cultured	Focioophile	Neutrophia	statator	Forthrocytes	CD4		CD8 CD14		CD19
Spink5	21	17	11			16	~~	15	28	36		104
chymase human	1221	47	65		45 1	12	101	62	108	59	104	45
tryptase alpha	21179	212	40		33 2	. 52		139	104	17	52	39
tryptase beta	25414	195	113		49 2	28	152	10	122	113	93	9
tryptase delta	349	45	Ū	6 2	23 5	. 22	74	113	=	10	42	Ξ
tryptase gamma	654	56	19		38 7	78	78	24	81	230	142	83
TRPV2	129	37	15		6 26	66	259	137	133	29	97	110
ANKTM1	28	28	~	8	38 3	30	96	18	14	46	Ξ	28
Cannabinoid receptor type I	20	41	14		47 3	36	27	19	41	26	18	54
Cannabinoid receptor type 2	160	369	226	5 578		177	271	530	324	232	212	421

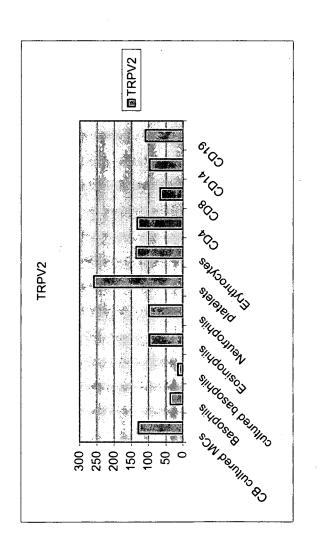


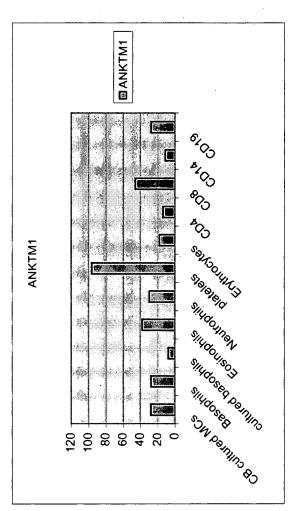


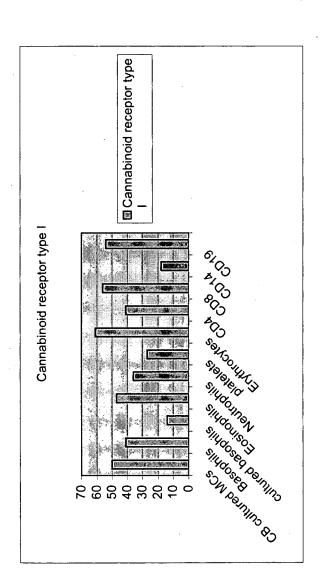


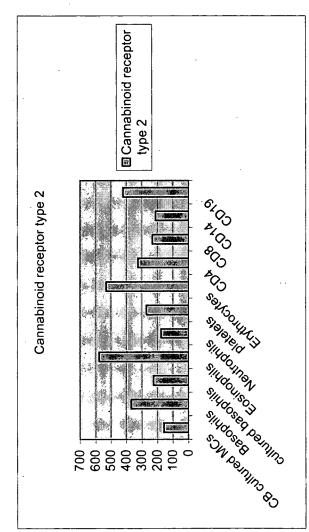




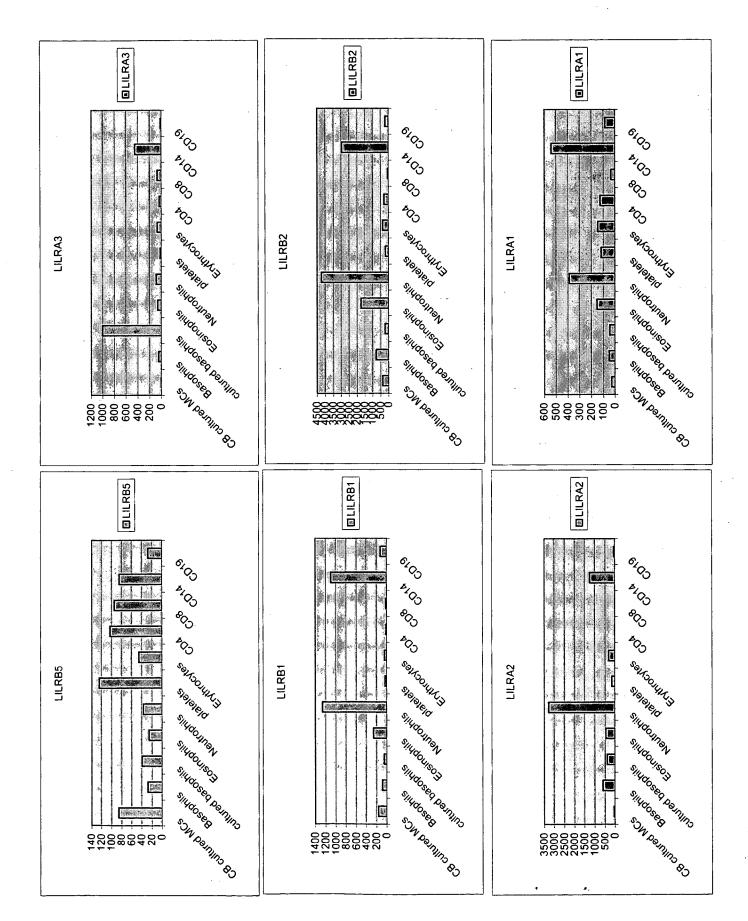


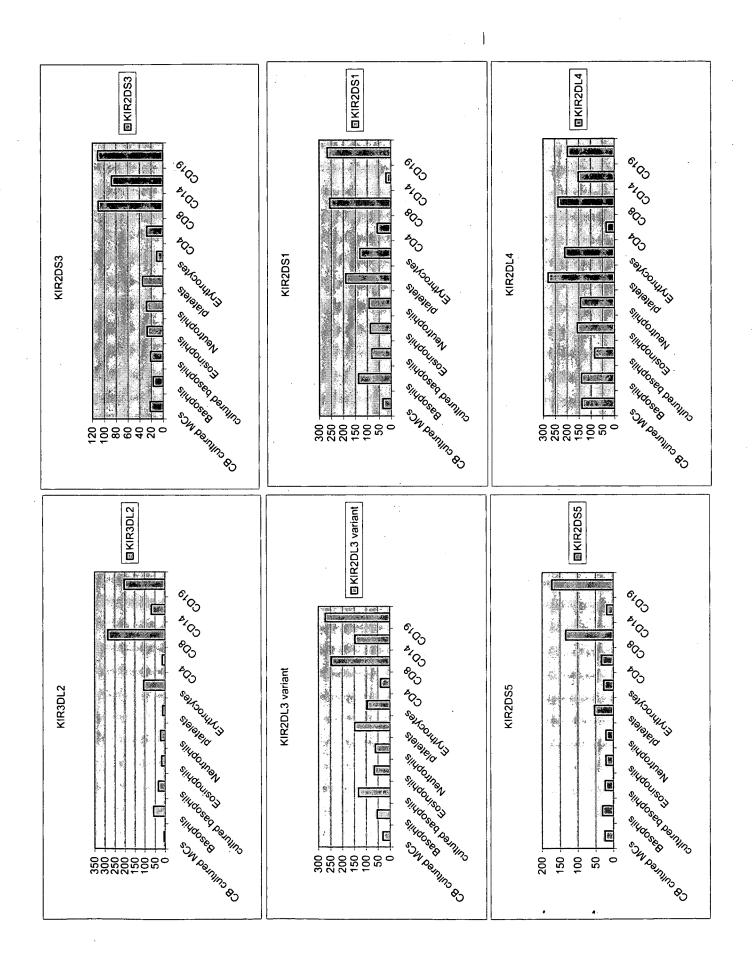






	CB cultured		cultured									
	MCs	Basaphils	basophils	Eosinophils	Neutrophils platelets	platelets	Erythrocytes CD4	es CD4	800	CD14	CD19	
LILRBS	98							46	103	95	82	28
LILRA3	7		_					89	31	89	451	13
LILRB1	165					31		42	13 ·	18	1105	125
LILRB2	375	5 780		_				375	586	58	2963	214
LILRA2*	46			1 461		146		318	8	6	1247	52
LILRA1	28							148	127	35	539	98
KIR3DL2	7	/ 58	34	1 18	23	13		106	14	286	89	204
KIRZDS3	22							Ξ	28	110	88	Ξ
KIR2DL3 variant	3							95	39	247	147	271
KIR2DS1	35							132	57	257	21	267
KIR2DS5	24							27	33	135	19	173
KIR2DL4	135				`			210	35	239	152	196
KIR2DL4								15	19	72	11	82
KIR-123FM	41							121	96	163	56	193
KIR2DL5.3	27							239	35	323	26	72
KIR3DL1	89							97	76	169	36	196
MIR cl-10	147	1 621		_				146	145	52	2154	66
PTPRF	82	. ,	4	5 28	37	&	9	12	13	14	32	23





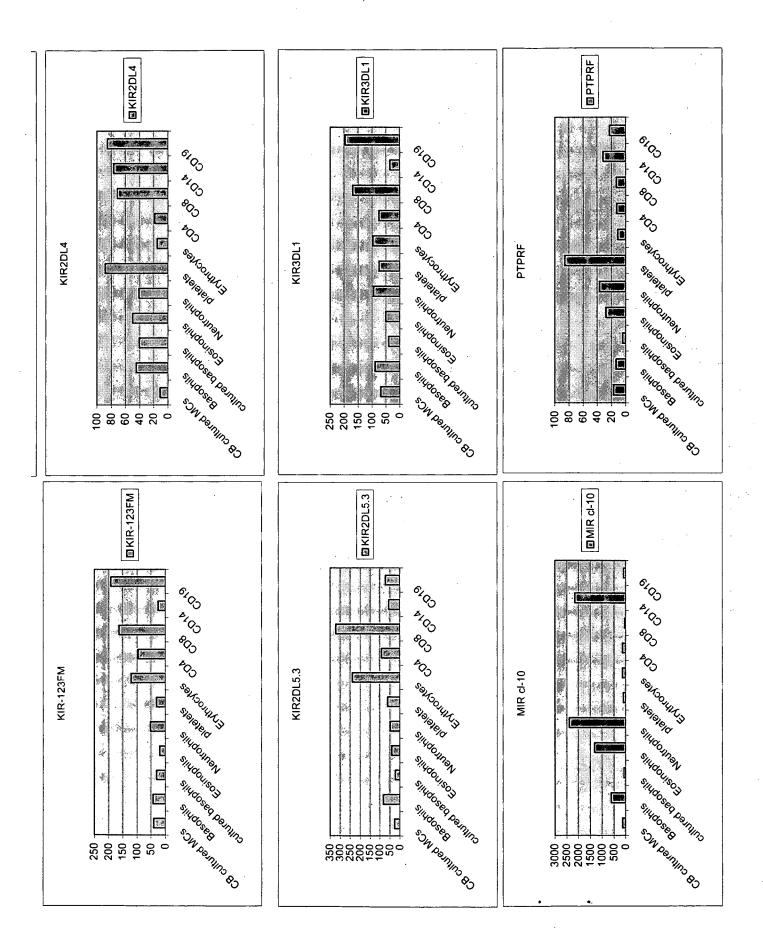


Figure 6A

Table E1. The complete list of granulocyte subtype-selective transcripts. Selectivity index (S.I.) was calculated by comparing the "normalized AD" level of a cell type or of two cell types with the maximal gene expression level of the other 8 or 9 cell types including platelets (PI), CD4⁺ cells, CD14⁺ cells, CD14⁺ cells, CD19⁺ cells and nasal polyp-derived cultured fibroblasts (Fb). When the result was accompanied by presence call, it was shown as a bold numeral. Italic numerals show that the raw AD levels were associated with absence call by the GeneChip analysis software. Transcripts having S.I. >3-fold were shown in A-H. Abbreviations used in the table through A-I were (small); the results obtained by the small sample protocol (see materials and methods), R; receptor, and ICN; ion channel.

A. Basophil (Ba)-selective transcripts (1/2).

				MC																			
Probe set	Accession #		Transcripts	cord	MC, fung	Ba 1	Ba 2 (small)	Ba 3 (small)	÷ €	2 Eo E	Eo 3 Er (small) (s	Eo 4 N (small) 1	Ne Ne	Ne 3 (small)	Ne 4 (small)	lg (ile	CD4	4 CD8	8 CD14	. CD19	19 Fb	æ	Ba S.1.
207539_s_at	NM_000589.1		11-4	0.2	0.1	10.4	16.9	13.4	0.1	0.0	0.2	0.1	2.1	3 0.0	٦	1.2 0.	2 0.0	0 0	2 0	1	0 0'	0 73	.348
210254_at	L35848.1	œ	HTm4	9.0	0.4	115.6	130.1	153.6	2.4	1.7	21.8	1.7	0.1 0	.9		.5 0	2 0.5	5 0.	4	1:	.2 0	0	38.24
205513_at	NM_001062.1		vitamin B12 binding protein	1.5	1.2	98.9	149.3	110.9	3.2	2.2	16.1	1.9	1.9	.5 6.3		1.1	8 0.7	7 0.	98	, A	0 6.	0.1 20	20.322
206148_at	NM_002183.1	œ	IL-3R	1.0	0.1	6.2	80.2	71.3	1.7	1.7	1.8	5.9	3.3	.4 0.2	~	.9 0.	2 0.3	3 0.	2 0		1.1		16.62
214920_at	R33964		FLJ11022 fis	0.1	17	4.9	13.1	15.4	0.1	0.2	0.7	9.5	1.0 0.	.4 0.5	2	0.7 0.	1 0.	1 0.	2 0		3 0	10	16.049
201825_s_at	AL572542		CGI-49	3.3	1.6	21.4	74.9	46.5	2.1	1.2	0.1	1.2	0.1	1,1 0.2	~	1.1 1.	3 0.7	7	7	A	.8 2	89 45	15.045
213238_at	A1478147		ATPase, Class V, type 10D	1.2	5.9	39.9	76.3	118.9	2.3	1.7	3.1	1.9	2.1	.4 2.	_	3.8 0.	7 0.9	9	9 2	εć.	4 0.	<u>+</u>	14.398
211734_s_at	BC005912.1	œ	Fc epsilon R Lalpha	10.5	28.4	210.0	220.8	226.2	8.4	2.1	8.4	5.9	3.3 0	7.	2	6.6 1.	4 5.1	.0	8	4.	0 5.	2 12	12.703
213894_at	BF447246		KIAA0960	0.1	0.0	3.7	13.0	15.6	0.3	0.4	9.0	0.4	0.9	.7 0.	_	0.6 0.	5 0.6	9	1 0	1	.2 0	2 12	12.272
206363_at	NM_005360.2		c-MAF	3.7	1.4	36.5	75.7	66.4	1.2	0.1	0.5	0.5	3.2 6	.1 0.	~	.0 0.	6 4.8	8 2.	7. 0.	0.5	0 0	0.0	11.927
203373_at	NM_003877.1		SOCS2	2.1	3.9	21.7	85.0	112.2	4.6	4.7	9.4	6.5	0.7	.2 0.	3	2.5	6 3.2	3.	.0	0.8	3.	9	9.8282
207538_at	NM_000589.1		114	0.3	0.1	5.4	8.6	9.5	0.4	0.4	0.0	0.3	0.1 0	.0 0.		0.7 0.	8 0.4	4 0.	3 0	0.1	0.0	6	9.8156
213684_s_at	BF671400		LIM-protein	9.0	0.1	1.1	21.0	9.02	6.1	1.6	5.4	8.	3.8	1.5		2.5 0.	1 0.0	0 0	3	0.	0.0	86	8.9245
209360_s_at	D43968.1		AML1b protein	10.4	2.4	53.2	131.1	90.5	8.5	7.0	13.8	11.3	0.5	.5 0.5		4.2 1.	3 4.1	1 5.	9	ω 	2.5 3	- 8	8.7543
220234_at	NM_004056.2		carbonic anhydrase VIII	0.4	0.1	11.7	10.2	9.0	1.2	0.1	0.5	0.4	0.8	1.0 0.1	_	3.5 0.5	5 1.1	1 0.	1 0		0.8 0	0.0	8.1309
210643_at	AF053712.1		osteoprotegerin ligand	0.1	0.3	1.8	3.4	6.0	0.7	0.4	0.3	0.5	0.4 0	1.1 0.3		1.0 1.0	1 0.3	3 0.	0.	5	0.2 0	0.2 7.1	7.6628
209211_at	AF132818.1		colon Kruppel-like factor	0.1	0.0	1.8	10.2	8.8	0.3	0.5	1.5	8.0	1.1 0	0.7 0.8	80	.5 0.	2 0.2	2 0.	9	?	0 1.0	3	7.239
204309_at	NM_000781.1		CYP11A	0.3	0.1	5.6	6.1	5.6	0.1	0.3	0.5	0.5	1.0	1.0 0.1	_	.9 0.	3 0.3	3 0.	1 0	0	.3	7.	6.8366
203372_s_at	AB004903.1		SOCS2	9.0	1.6	9.5	8.5	15.9	1.4	2.4	6.0	0.5	0.7.0	1.6 0.4		.8 0.	9.1.6	6 1.	3	3	1.3	1.4	6.8271
207463_x_al	NM_002771.1		serine protease 3 (trypsin 3)	1.0	1.0	8.8	10.0	10.9	0.4	1.0	0.	1.2	1.2 0	.9 1.	6	.8	4 0.6	0 9	9	9	7.1	.2	6.7218
213624_at	AA873600		acid springomyelinase-like phosphodiesterase	4.7	2.0	20.8	30.6	25.9	2.1	2.8	3.8	1.7	1.5	1.1		.3 0.	1 0.2	2	4	8	0.2 3	6	6.6846
214873_al	AL137651.1		done DKFZp43400213	0.1	0.4	3.7	15.5	24.4	5.0	6.0	2.5	2.3	0.2	1.1 0.7	_	0.9.0	1 1.	1	5 0	.5	0 9:0	7	6.2585
204928_s_at	NM_019848.2		protein P3	2.3	1.5	8.6	49.0	34.3	3.7	3.3	8.8	4.2	1.6	.1 0.	•	2.2 4.	1.1	3.	0	8	2 0.3	9	6.2142
208935_s_at	L78132.1		prostate carcinoma tumor antigen (pcta-1)	2.2	1.2	6.6	19.1	16.6	2.3	1.5	3.6	2.5	-	7	•	0.9	8 1.8	8 2.	0 2	O	1.4	9	6.2011
203201_at	NM_000303.1		phosphomannomutase 2 (PMM2)	1.8	0.3	10.7	15.9	6.5	6.0	9.0	1.2	6.0	0.8	1,4	1	.3	0 1.0	. .	2				6.1557
201826_s_at	NM_016002.1		CG1-49	1.8	3.1	11.0	27.5	15.1	1.6	1.1	4.4	2.1	1.1	.t.	3	1.8 1.	6 1.		1.1	· ·	0.6 2	2.9 5.	9407
213421_x_at	AW007273		serine protease 4 (trypsin 4)	1.8	1.7	5.9	12.5	11.4	1.0	0.5	1.2	0.5	1.2	7.	~	.2 0.	9 1.0	7.	2 +	0.	14 1	5.	4802
209348_s_at	AF055376.1		c-MAF, short form	6.4	4.9	29.6	47.7	42.3	8.0	0.3	0.4	9.0	9.3	0 0	6	1.	0 7.3	33	-	6.0	1.7	4.	5.3183
213343_s_at	AL041124		hypothetical protein PP1665	0.9	0.2	12.2	17.0	12.0	3.5	1.7	0.5	9.0	0.1	.3 0.8	8	0.4	6 1.5	5 2.	0 9		.6	5	2721
202491_s_at	NM_003640.1		I kappa B-associated protein	1.6	3.8	23.9	51.9	43.5	4.6	4.0	1.1	7.3	1.6	.1	10	2.3	7 3.7	7.	£.	, ,	1.7	رب دن	5.107
221021_s_at	NM_030877.1		Bos taurus P14 protein	6.9	3.2	7.2	29.2	58.9	2.8	3.2	5.4	6.3	7.9	.2	80		6 2.3	7	7 .	e.	1.5	86 4	.9129
213346_at	BE748563		hypothetical protein BC015148	2.7	6 .	17.5	38.8	25.8	0.4	6.7	6.2	8.8	1.4	.0	6	1.1 0.	2 1.	2 1.	5 0	9.	-	ون <u>ج</u>	4.8879
209764_at	AL022312		mannosyl (beta-1,4-)-glycoprotein beta-1,4-N-acetylglucosaminyltransferase	0.0	0.9	4.1	6.5	5.8	0.8	1.4	6.0	4.4	0.6	.0	10	0.2 1.	0	0.	9	.2	0 9.		4.855
207067_s_at	NM_002112.1		histidine decarboxylase	64.1	14.7	105.2	164.9	165.7	3.4	1.4	1.4	2.8	2.1	.2 4.6	•	14.0 2.	4 0.8	9	9 1	.1	0 7.0	0.2 4.	4.6305
210375_at	X83858.1	GPR	prostaglandin E receptor, type 3a2	1.0	9.0	2.4	11.5	17.0	0.3	0.1	0.1	0.0	0.4 0	0.7 0.9	6	0.7 0.	6 0.3	3 0.	2 0	0.2 (1.5	1.7 4.	4.6103
206306_at	NM_001036.1		ryanodine receptor 3 (RYR3)	2.1	1.1	3.7	11.0	7.8	1.0	9.0	1.5	9.0	1.2	.3 1.	ıc.	.9 1.	1 0.7	7 0.	٤.	0.1	0.2.0	4.	4.5552
210001_s_at	AB005043.1		SOCS1	5.2	0.2	3.0	24.1	29.9	2.1	3.0	3.2	3.4	0.3	.0	6	.3 0.	8 0.1	9	9	0.0	9.8	<u>6</u> ,	.5248
20,4614	1 37 3 COO MIN		serine (or cysteine) proteinase inhibitor, clade B (ovalbumin), member 2	,	9		120	12.5		70	2	5	9	40			40		,	,	,		4 5061
200060 at	V16272 1		honotondo grouph factor (UGE)	3 6	2 0	9 6	; ;		, u			, ,									, ,	100	4 4335
209212 s at	AB030824.1		transcription factor BTEB2	0.4	0.5	7.5	14.3	22.1	. 62	2.0	6. 6.	3.7	1.2	1.3 1.5		2.5			0.2	: :	•	_	4.363
													l	l				l				ł	١

A. Basophil (Ba)-selective transcripts (2/2).

Figure 6 B

Probe set Acc	Accession #	Transcripts	cord	MC. fung	Ba 1	(small)	(small)	3 -	2 ((small) ((small)	2 -	2 ((small)	(small)	Į,	CD4	CD8	CD14 (CD 19	۽ ۾
214651 s. at U41	U41813.1	class I homeoprotein (HOXA9)	9:0 ·	9.0	1.7	6.3	8.3	0.3	0.5	8.0	0.5	0.1	0.1	0.5	1.0	1.0	0.1	0.1	0.3	0.1	0.1
	AL041124	DKFZp434D0316_s1	1.6	1.9	14.6	22.5	17.4	4.0	2.1	0.7	1.4	1.1	1.0	9.0	0.4	2.3	2.5	4.1	2.5	2.3	8:
		catenin (cadherin-associated protein).	:		;	;	;	;			;	,	;	;	;	,	;	;	;	,	ç
	NM_004389.1	alpha 2	0.5	0.1	⊋ .	3.9	3.5	9.0	0.1	0.1	0.1	0.1	0.1	0.5	0.1	0.1	0.1	9.0	0.2	0.2	0.2
	NM_025231.1	FLJ22191	0.3	0.7	2.5	, S	12.0	0.9	0.5	5.0	÷.	0.2	1.1	4.2	2.2	9.0	7.	4.	0.3	4.	0.
	AK026415.1	beta2-chimaenin	4.7	. .	8.3	22.9	21.8	-	9.0	6.0	1.0	0.5	1.5	0.8	1.0	0.1	6.0	9.	3.3	0.5	0.4
214637_at BG4	BG437034	oncostatin M	0.4	0.0	6.0	5.4	4.4	0.2	0.7	0.1	0.7	0.5	1.8	0.2	1.1	0.4	0.1	0.1	0.5	0.1	0.0
202490_at AF1	AF153419.2	I kappa B-associated protein	0.2	0.2	1.3	3.6	2.2	0.4	0.4	0.5	0.1	0.3	0.0	0.2	0.0	0.3	0.3	0.5	0.2	0.4	0.2
	NM_024837.1	FLJ21472	4.1	1.5	9.1	9.7	12.2	1.9	3.3	3.0	1.9	9.0	9.0	Ξ	6.0	0.7	9.0	0.5	9.0	0.3	0.0
218318_s_at NM	NM_016231.1	nemo-like kinase	1,4	1.3	6.2	7.0	7.1	9.0	6.0	8.0	9.0	0.5	1.0	0.1	0.1	1.6	1.3	6.0	1.2	Ξ	4.0
	AW166925	FLJ14135 fis, clone MAMMA1002728	0.1	0.1	2.1	3.5	5.	0.5	0.7	8.0	6.0	0.0	0.1	0.1	0.1	0.5	0.3	8.0	0.4	0.5	0.1
	AV700891	ETS2 intronic transcript 1 mRNA	9.0	1.4	31.1	86.8	63.8	3.6	2.2	5.9	3.5	10.8	10.5	. 16.6	19.0	1.2	0.7	0.7	1.8	0.3	9.0
	AL136877.1	DKFZp434F205	2.8	5.0	17.3	27.7	31.8	4.4	5.3	9.1	8.3	4.3	3.9	5.7	5.5	0.1	4.8	4.2	2.5	4.1	4.8
ŧ	AB002356.1	MAP-kinase activating death domain	9.6	4.6	12.7	35.0	40.7	6.2	6.1	7.0	8.3	2.2	1.3	1.9	3.3	5.5	2.7	5.4	3.0	3.5	1.1
		v-ets avian erythroblastosis virus E26																			
	AL575509	oncogene homolog 2	1,	0.1	18.4	34.7	20.7	2.8	1.6	1.7	8 9.	6.4	6.6	8.7	8.9	0.5	0.9	1.0	4.4	0.1	-:
a a	NM_022754.1	FLJ12876	1.2	1.0	3.5	7.5	7.4	1.4	1.0	2.7		1.0	9.0	6.0	0.1	6.0	4.4	0.4	0.7	0.5	4.
	NM_001813.1	FLJ14150 fis. clone MAMMA1003026	0.0	0.3	1.5	8 .	13.1	1.7	0.5	6	1.2	0.5	1.3	.	6.0	-	.	6.0	1.0	9.0	0.5
	AF312230.1 GPR	histamine H4 receptor	9.0	0.8	35.9	34.2	32.5	7.8	6.1	8.7	9.3	0.5	9.0	8.0	1.0	0.4	8.0	0.5	9.0	0.0	0.7
201663_s_at NM	NM_005496.1	chromosome-associated polypeptide C	2.2	4.3	17.6	15.1	18.1	3.4	9.9	4.9	4.2	4.0	3.7	2.2	2.7	1.7	4.	5.9	1.7	3.7	4.2
208933 s at Al6	A1659005	fectin, galactoside-binding, soluble, 8 (natectin 8)	11.8	6.9	26.7	28.3	28.2	6	9	10.1	8.	80	5.	40 40	2.1	6.3	9.7	7.4	7.2	6.	2.7
	AL563460	GATA-binding protein 2	50.4	38.9	141.1	189.2	154.7	4.4	£.	3.8	3.1	0.7	1.2	5.4	10.8	2.5	0.7	6.0	0.3	0.1	6.3
		fatty-acid-Coenzyme A ligase, very	:		;	,	,	,			•	,	,		ć	Ş	;	į	ò	6	
	NM_003645.1	long-chain 1 (FACVL1)	£	2.5	0.7	16.2	3.6	0.3	. O. 3	6.0	c.5	0.1	2.	0.2	5.5	7.	6.0			ۍ . د د	<u>.</u>
_	D86962.1	KIAA0207	9. 7.	5.6	12.5	38.0	33.1	6.9	4.2	12.3	10.4	1.7	1.7	2.1	æ.	4.	0.4	0.7	. .	0.4	8. 6
	AB002356	KIAA0358	2.5	4.	13.8	24.6	29.8	.	6.4	9.6	7.0	3.0	3.2	E. :	4		3.3	5.3	3.7	4.4	2.5
	NM_003914.1	cyclin A1	5.6	9.0	16.2	20.6	10.5	0.1	0.6	6.0	0.1	9.0	0.1	0.8	0.4	1.9	0.7	0.1	0.1	0.3	8.0
	NM_012097.1	ALJP-nbosylation factor-like 5	2. 2	7	37.5	9.5	9.4	- c		5.4	ניאַר .	, .	0.0		o •	ים פים	, c			, c	
	Al33883/	zuotin related factor 1	; ;		2 0 (5.5.	, o .	5 . 0		4, 4 10, 6		٠	7.6	? :	٠,	5 6	4 6	· ·	3 ;		9 9
208158_s_at NM	NM_018030.1	oxysterof-binding protein-related protein	1.2	2.3	89	18.0	15.3	9.0	8.0	9	4.	.	2.3	2.7	8 .	5.0	0.2	0.7	.	0.5	4.0
210109 at 7 AF1	AF 191492.1	nasopharyngeal carcinoma associated gene profein-8	1.2	0.7	3.6	6.4	4.2	1.2	9:	1.0	0.3	9.0	1.1	1.7	9.0	0.2	1,3	0.5	4.	9.0	0.1
	NM 025143.1	FLJ20856	1.0	1.1	26.1	37.3	26.2	6.2	6.5	14.1	11.3	3.3	2.5	2.0	2.0	1.1	6.0	6.0	3.5	1.1	0.1
	, 00170	aml 1 (acute myeloid leukemia 1)	,	ċ	;	,	,		4	4	;	ć		4	;	,		,	à	90	-
	1330.1	oncogene	7.		- c	4 6	: ;	9 9		? ;	- 6	3 6		5 .		5 6	3	2 5	9 6	3	; ;
5	NM_030941.1	exonuclease NET-sp	5.5	9.9	9.	30.5		0.	5.3	7 ;	8.5	5	7.0	9	9.5	0.7		?; •	7.0	÷ ;	7.0
212412_at AV/	AV/15/6/	URF-Zp564A072 chromosome 16 BAC close	18.7	7.0.	5.	C.7.	5.70	:	:	20.5	e E	ė.	6.5	- n		ò	-	-	9.0		3
215215 s at ACC	AC004381	CIT987SK-44M2	2.9	2.5	4.8	12.8	10.4	9.0	6.0	1.4	0.8	0.1	0.0	0.2	9.0	0.1	0.4	0.3	9.0	0.3	0.5
	AB014731.1	SMAP-3	9.7	5.4	13.2	48.5	32.4	5.2	5.7	18.1	10.7	3.5	3.8	11.4	8. 1.	8.2	5.6	6.0	4.1	7.5	8.0
	NM_018439.1	hypothetical protein IMPACT	1.3	2.4	5.9	10.7	9.3	5.5	1.5	2.7	2.1	9.0	0.7	9.0	9.0	1.2	8.0	0.7	6.0	Ξ	7.
	NM 018191.1	hypothetical protein FLJ10716	0.8	0.5	6.4	13.7	8.7	2.3	1.7	3.4	3.9	0.1	5.0	2.8	3.6	0.8	2.3	5.0	1.7	2.2	2.4
	AI081194	KIAA0379	2.1	3.8	1.1	26.7	22.4	2.5	1.5	3.4	2.5	1.9	1.1	2.3	9.1	4.6	2.5	1.3	0.1	2.2	6.1
ä	D89788.1	aml 1 (acute myeloid leukemia 1) oncogene	1.5	0.5	3.1	9.4	7.5	2.0	2.5	1.7	1.7	1.0	1.4	0.3	1.1	0.4	0.4	9.0	1.1	0.7	0.2
	426406	lectin, galactoside-binding, soluble, 8	;	;	•	č	7 9	•	•			,	;	2.2	4		•	,			0
	136103	(galecuii o)	; ;		,				o e	9 9		· •		; ;			<u> </u>	<u>.</u> .	: ;		9 6
203164_at BE4	BE464756	acetyl-Coenzyme A transporter	6.3	5.	3.4	12.6	17.4	6.7		9 .0	7.7		o.	7.	?	.	?	0.7	- •	;	Ç.
205768 s at NM	NM 003645.1	fatty-acid-Coenzyme A ligase, very Iong-chain 1 (FACVI.1)	3.0	1.7	3.9	10.7	7.7	0.4	0.1	9.0	0.5	0.1	9.0	0.1	0.7	0.0	9.0	0.1	0.2	0.1	0.1
	AB003476.1	A kinase (PRKA) anchor protein (gravin) 12	25.1	6.4	44.0	69.5	59.6	1.5	0.4	9.0	0.8	0.1	0.5	1.0	3.0	6.0	0.8	0.5	0.1	0.3	18.8
		Ca2+-independent phospholipase A2 short									•										

Probe set	Accession #		Transcripts	MC, blood	MC.	- Ba	Ba 2 (small)	Ba 3 (small)	Eo 1	Eo 2	Eo 3 (small)	Eo 4 (small)	S -	2 N	Ne 3 (small)	Ne 4 (small)	· 1	CD	CD8	CD14	CD19	. و	Eo S.I.
207328 at	NM 001140.1		15-lipoxygenase	0.1	9.0	0.7	0.1	0.1	14.5	24.6	18.3	17.0	0.1	1.1	0.1	0.1	9.	0.0	0.1	0.0	0.1	0.0	74.129
219695_at	NM 024703.1		FLJ22593	0.0	1.0	6 .	8.	-	29.8	34.7	24.8	27.8	0.8	1.5	0.4	6.0	0.1	9.0	0.4	0.3	0.1	0.1	19.123
208253_at	NM 014442.1	α	sialic acid binding Ig-like lectin, Siglec 8	1.3	2.4	0.4	0.4	0.5	17.1	23.1	17.6	11.8	0.1	0.4	0.1	0.8	0.3	0.2	0.1	0.2	0.5	0.2	9.8056
211922_s_at	AY028632.1		catalase	3.7	2.2	6.2	8.9	2.2	78.2	134.5	119.7	88.8	16.2	13.0	8.8	8.7	0.7	2.2	5.9	1.1	7.5	4.6	9.125
201802_at	NM_004955.1		solute carrier family 29 (nucleosidetransporters)	4.8	2.8	5.9	3.4	1.9	24.8	30.4	36.1	42.3	0.1	9.0	9.0	0.8	0.5	6.0	0.3	2.2	1.0	1.1	8.989
214523_at	NM_001805.1		CCAATenhancer binding protein (CEBP), epsilon	0.1	0.1	0.7	1.2	1.2	7.9	11.4	23.4	17.3	1.8	1.5	6.1	1.1	4.4	0.5	9.4	1.0	0.1	0.1	8.9462
210029 at	M34455.1		interferon-gamma-inducible indoleamine 2.3-dioxygenase	0.7	0.1	53	4.2	5.6	20.0	39.0	33.4	29.5	1.6	2.1	2.3	1.8	1.7	1.5	1.7	0.3	4.4	0.9	7.7078
215573_at	AU147084		FLJ12072	0.1	0.1	0.5	0.3	1.1	8.6	8.4	16.9	7.2	0.4	1.9	1.8	2.0	0.1	0.1	0.0	0.1	0.4	0.0	7.5983
201801_s_at	AF079117.1		sotute carrier family 29 (nucleosidetransporters)	3.5	6.	1.7	0.5	6.0	29.7	47.1	7.4	11.0	0.1	0.5	0.4	0.7	1.1	9.4	0.1	4.	9.0	0.1	7.0314
213825_at	AF221520.1		oligodendrocyte lineage transcription factor 2	0.4	9.0	0.5	0.4	9.0	6.0	10.9	18.7	12.5	9.0	6.0	1.5	1.0	9.0	1.7	0.3	0.4	0.3	0.3	6.5987
219821 s at	NM 018988.1		glucose-fructose oxidoreductase domain containing	3.0	4.	4 .8	2.6	2.3	18.7	17.3	30.9	31.3	3.9	3.1	2.4	2.9	3.2	1.3	2.5	3.0	3.8	0.8	6.2284
205472_s_at	NM_004392.1		dachshund (Drosophila) homolog	0.0	0.1	. 0.1	0.4	0.1	2.7	1,4	2.2	1.7	0.3	0.1	0.1	0.1	0.5	0.1	0.1	0.1	0.3	0.0	6.0088
202188_at	NM_014669.1		KIAA0095	.0.5	6.0	0.4	1.8	1.2	6.8	8.5	13.5	10.6	0.5	0.2	0.2	0.1	0.2	0.3	1.7	1.7	4.4	0.4	5.4499
210548_at	U58913.1		chemokine CCL23	9.0	1.8	9.0	1.1	0.5	5.5	6.2	4.2	6.9	. 0.1	0.9	0.1	0.2	0.3	0.1	0.5	9.0	0.1	0.0	5.3082
209447_at	AF043290.1		fyriphocyte memorane associated protein (8B7)	9.0	0.5	2.5	3.4	3.6	41.8	49.7	66.2	60.1	0.5	6.0	0.5	1.2	5.6	5.1	10.4	1.6	6.3	2.7	5.1472
206171_at	NM_000677.2	GPR		2.3	8.2	8.7	3.7	2.2	7.8	13.9	22.0	17.9	1.6	3.1	2.2	2.5	1.2	1.6	0.5	2.0	0.5	0.5	5.0469
210549_s_at	U58913.1		chemokine CCL23	1.1	1.1	0.1	1.3	9.0	7.9	5.6	2.7	8.4	0.0	9.0	0.0	0.2	0.1	0.1	0.1	0.2	0.0	0.4	4.9873
214183_s_at	X91817.1		transketolase-like protein	0.1	0.2	0.5	0.1	0.1	3.2	4.0	9.9	7.2	0.5	0.5	0.1	0.1	0.5	0.5	1.0	0.1	9.0	0.0	4.7894
215350_at	AB033088.1		envelope 1	0.1	0.1	0.5	0.1	0.1	3.8	4.2	3.0	2.8	0.0	0.1	0.2	0.4	0.7	0.2	0.1	0.0	0.1	0.5	4.7822
206277_at	NM_002564.1	GPR	P2Y2 purinergic receptor	0.1	0.1	0.1	0.5	0.1	3.4	4.7	8.1	5.9	0.1	0.3	0.1	0.1	0.5	0.1	0.3	1.2	0.2	0.1	4.3043
204776_at	NM_003248.1		thrombospondin 4 (THBS4)	0.8	0.1	1.5	0.1	0.8	3.2	2.9	10.0	6.4	1.8	4.4	9.0	6.0	03	0.5	0.3	1.4	1.1	1.2	4.1746
201563_at	L29008.1		L-iditol-2 dehydrogenase	7.	5.6	3.5	7.4	12.8	22.5	59.9	49.0	37.0	6 .	3.0	4.4	2.2	~	9	7.	2.7	5.5	-	4.1557
206637_at	NM_014879.1	GPR	P2YX purinergic receptor GPR105 for UDP-glucose	5.7	1.6	13.3	19.9	13.5	44.2	55.1	81.4	62.4	1.8	9.3	15.4	9.5	0.7	6.0	9.0	0.0	5.5	0.1	3.8761
213622_at	AI733465		collagen, type IX, alpha 2	1.5	1.4	6.	1.2	1,3	10.3	9.6	9.9	8.8	2.7	2.1	2.1	2.2	0.7	7.5	1.2	2.2	7.	6.0	3.8576
214705_at	AJ001306.1		PDZ domain protein	0.1	0.1	0.4	0.3	0.1	2.1	1.5	5.6	2.2	0.3	0.0	9.0	0.7	0.1	0.4	0.3	0.3	9.0	0.1	3.6532
266_s_at	L33930		CD24 signal transducer	0.5	0.1	e ;	0.3	8. 6	11.9	14.0	18.8	13.9	ō ;	0.5	0.3	0.8	6.0	0.0	0.1	0.1	4 6	0.5	3.3793
201432_all	1.26/100 MM		caraiase (CAT)	2.5	5.4	7.00	4 4	5.0	602.3	23.7	0.122	5.702	9 6	9 . u	3 5	6.00	4. 6	9 6		9 4			3 3080
209090_ai	AK000168 1		CD24 signal transducer		4 4	16.2	6.0		66.1	70.1	81.9	77.9	1.7	2.8	5.6	4	8.	0.5	0.6	6.0	22.5	0.5	3.2741
			lysosome-associated membrane		. :				;	;	,	;	,	;	:	;	;		į	;		,	
205569_at	NM_014398.1		glycoprotein (TSC403)	0.4	0.2	.0.	0.1	0.4	2.4		e ;	D (9.0		4.	, c	3 6	2 ;	`; ;	ŝ	0.0	2 6	3.2308
219233_s_at	NM_018530.1		hypothetical protein PRO2521	0.1	1.9	2.1	4.6	5.4	7.5	8.5	21.8	21.2	0.8	0.1	1.5	2.1	0.1	1.1	4.	0.1	4.	0.5	3.2093
202286_s_at	304152		gastrointestinal tumor-associated antigen GA733-1	0.9	0.7	1.3	0.3	0.1	7	2.5	8.7	1.8	0.4	0.8	9.0	9.0	0.5	0.2	0.7	0.3	0.5	0.1	3.1844
206442_at	NM_003007.1		semenogelin I (SEMG1)	9.0	0.1	0.4	0.0	0.1	1.2	3.2	4.9	3.4	0.3	0.5	1.1	1.9	0.1	0.2	0.2	0.0	0.4	0.1	3.1837
205733_at	NM_000057.1		Bloom syndrome	1.9	1.1	5.0	2.4	2.2	8.5	6.0	7.5	11.2	2.1	2.0	1.0	1.2	1.2	2.4	1.8	1.1	2.6	1.2	3.1427
204392 at	NM 003656.2		calciumcalmodulin-dependent protein kinase I (CAMK1)	4.0	1.1	6.3	6.8	6.4	17.8	19.3	24.9	26.5	9.0	0.2	. 4.8	1.5	0.1	4.4	1.0	3.0	9.0	6 .	3.0763
213497_at	AL050374.1		DKFZp586C1619	1.0	0.7	1.0	1.3	1.6	6.3	6.8	6.8	6.0	2.3	3.0	1.4	2.3	2.0	0.4	0.5	1.6	1.1	0.8	3.0099
219296 at	NM 019028 1		similar to ankyrinrepeat-containing	2.2	1.7	00	60	9,	7.3	4.4	4.8	7.4	8	2.4	2.4	2.3	0.1	1.4	1.6	1.3	6.1	8.0	3.0018
£10000 00						ţ									i								

C. Neutrophil (Ne)-selective transcripts (1/7).

Probe set	Accession #		Transcripts	MC. cord blood	MC, lung	Ba 1	Ba 2 (small)	Ba 3 (small)	Eo 1	Eo 2 (8	Eo 3 E (small) (s	Eo 4 (small) N	Ne 1	Ne 2 (s	Ne 3 N	Ne 4 (small)	D ld	CD4 C	СОВ	CD14 C	CD19 F	۾	Ne S.I.
205403_at	NM_004633.1	œ	interleukin 1 R. type II	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1		37.4	41.4	73.8	61.5	0.2	0.4	0.1	0.2	0.0	0.0	127.86
216782_at	AK026679.1		FLJ23026 fis	0.0	0.2	-0.5	0.1	0.1	0.1	0.1	0.5	0.2	33.2	31.5	33.0	17.4	0.1	0.1	0.1	0.1	0.2	0.2	112.28
210119_at	U73191.1	C	channel Kirt.3	0.7	1.1	0.1	0.1	0.5	0.1	0.1	6:0	9.0	81.6	98.6	129.0	88.8	0.7	0.2	0.0	8.0	0.0	0.0	107.47
209395_at	M80927.1		glycoprotein-39)	1.5	0.2	6.0	0.1	0.4	0.3	0.1	1.0	0.1	28.9	25.3	7.07	40.7	0.1	0.0	0.1	0.1	0.1	0.0	79.595
203691_at	NM_002638.1		professe frimpilor 3, Skin-derived (SKALP)	0.2	0.1	0.1	0.1	0.8	0.5	0.2	9.0	0.1	16.9	36.7	27.7	24.5	0.1	0.1	0.1	0.1	0.5	0.1	51.901
211372_s_at	U64094.1	œ	interleukin 1 R. type II	0.1	0.2	0.1	0.1	0.0	0.1	0.5	0.7	0.1	21.6	36.5	31.4	29.9	0.3	0.2	0.0	9.0	0.0	0:0	. 909:05
207008_at	NM_001557.1	GPR	CXCR2 interleukin 8 receptor, beta	0.0	9.0	Ξ	1.2	1.0	1.3	1.3	2.1	1.2	129.8	168.4	5.18	8.89	2.7	0.3	8.0	0.7	9.0	0.0	39.316
206515_at	NM_000896.1	1	(CYP4F3)	0.3	0.1	Ţ.	0.7	1.2	0.5	9.0	5.9	1.7		40.3	57.2	48.9	0.5	6.0	0.0	0.7	0.3	0.3	34.919
204007_at 204470_at	J04162.1 NM_001511.1	œ	Fc gamma R IIIb (CD16) melanoma growth stimulating activity, alpha	1.2	0.0	7.0	9.9	1.8	1.1	1.8 0.5	2.8	1.9	204.5 19.1	226.5	194.0 26.3	173.3 48.1	6.6 0.1	1.3	1.3	2 1 2	2.6	0.7	29.895 28.189
206025_s_at	AW188198		tumor necrosis factor. alpha-induced protein 6	0.1	0.3	0.1	9.0	9.0	9.0	0.1	0.2	0.1	19.0	29.9	17.2	24.7	0.1	0.2	0.2	8.0	0.3	9.0	26.336
209396_s_at	M80927.1		chitinase 3-like 1 (cartilage glycoprotein-39)	1.8	0.2	9.1	0.4	0.5	8.0	0.5	5.6	6.0	37.9	32.2	52.8	26.7	1.4	0.0	0.0	0.1	0.1	0.4	25.669
211806_s_at	D87291.1	Š	Charles inward rectiner potassium channel Kirt.3	0.9	1.7	2.0	1.4	1.4	2.1	1.3	1.8	1.7	62.2	77.2	77.5	26.8	3.0	1.0	1.1	1.7	6.0	0.7	22.254
221920_s_at	BE677761		mitochondrial solute carrier	0.4	0.8	6.9	0.0	0.2	2.4	5.5	2.2	1.5	29.0	43.2	57.4	46.8	2.5	0.7	0.3	8.	<u>-</u> :	0.	20.163
207094_at	NM_000634.1	GPR	atpha	0.3	0.1	4.2	4.3	4.4	0.3	6.0	0.2	0.3	69.2	81.9	95.3	87.4	0.4	0.3	0.2	0.5	0.1	0.1	19.325
213589_s_at	AW468201		23614 mRNA sequence	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.5	0.1	5.5	2.7	5.1	4.0	0.1	0.1	0.2	0.1	0.1	0.0	18.973
218963_s_at	NM_015515.1		DKFZP434G032 PAR2 proteinase activated	0.0	0.1	0.5	0.3	0.1	Ξ	2.8	8.2	2.0	46.3	32.5	40.3	31.3	0.3	0.3	0.1	0.1	0.1	0.1	18.3
213506_at	BE965369	GPR	receptor-2	0.1	0.0	0.0	0.4	0.3	1.2	1.3	4.1	2.3	33.4	35.7	45.4	33.4	0.1	0.7	0.1	1.6	9.4	7.7	18.208
220187_at	NM_024636.1		FLJ23153	0.3	0.1	0.3	0.1	0.1	0.0	0.1	0.5	0.5	16.5	26.3	5.7	7.0	0.0	0.1	0.1	0.7	0.1	0.2	17,111
206026_s_at	NM_007115.1		alpha-induced protein 6	0.1	8.0	9.0	9.0	9.0	Ξ	0.5	6.0	0.1	18.2	23.3	20.0	17.0	9.0	1.0	1.0	7	0.1	<u>0</u> .	17.051
41469_at	L10343		elafin	1.2	0.7	1.2	9.0	9.0	6.0	1.0	0.7	9.0	16.2	39.1	19.5	17.4	1.3	1.4	0.7	9.0	0.5	0.5	15.913
205568_at 210483_at	NM_020980.2 BC005043.1	S C	aquaporin 9 decoy receptor 1, TRAILR3	0.5	0.9	0.0	0.0	0.1	1.0	0.9	0. t .	1.2	33.3 25.8	136.7 23.2	169.0 9.7	110.5 8.6	0.7	0.9	0.3	8.6 0.2	0.7	0.7	15.805
215223_s_at	W46388		superoxide dismutase 2	4.4	1.9	1.7	4.9	8.7	5.8	3.3	6.0	8.3	124.6	153.1	142.4	151.8	11.0	9.1	77	7.3	2.2	1.5	13.008
210484_s_at	BC005043.1	α	decoy receptor 1, TRAILR3	0.9	6.0	6.0	0.5	0.1	2.2	4.8	9.0	9.0	60.1	86.3	4.7	5.9	0.5	9.0	0.4	6.0	0.4	0.5	12.748
205654_at	NM_000715.1		protein, alpha	0.5	1.1	1.3	0.7	0.7	1.5	1.0	1.3	1.4	2.1	20.7	43.6	33.7	1.2	0.3	1.3	1.2	1.2	6.0	12.161
210773_s_at	U81501.1	GPR	formyl peptide receptor 2	0.8	0.1	9.0	9.0	0.3	9.0	9.0	9.0	0.5	86.9	105.0	20.5	59.9	0.1	1.0	9.0	6.0	0.8	0.1	12.14
206222_at	NM_003841.1	α	decoy receptor 1, TRAILR3	1.5	9.0	6.	0.7	0.7	3.6	13.7	3.4	3.9	21.0	137.1	37.5	22.5	0.0	0.1	0.1	8: ;	0.5	7. 0	12.034
202083_s_at	NM_003003.1		SEC14 (S. cerevisiae)-like 1	8.0	0.2	6.0	œ.	3.0	2.2	5.6	3.8 8.	5.6	36.7	25.9	33.9	32.6	 	0.1	r.0	5	9.		67/:11
211163_s_at	AF012536.1	œ	decoy receptor 1, TRAILR3	0.2	0.1	2.2	1.5	9.0	5.6	12.3	9.4	6.4	87.9	110.9	60.4	55.8	0.5	0.2	0.1	0.7	0.1	9.0	11,404
205931_s_at	NM_004904.1		protein CRE-BPa	0.0	0.1	0.3	0.1	9.0	9.0	9.0	9.0	0.7	23.2	14.5	20.5	12.9	9.1	9.0	0.1	9.	0.1	0.1	10.618
205922_at	NM_004665.1		vanin 2	0.1	0.0	1.2	5.6	3.9	1.0	0.2	1.9	0.3 1	14.5	114.6	174.1	189.2	0.1	1.4	4.0	13.8	1.7	0.0	10.451
210176_at	AL050262.1	œ	Toll-like receptor 1	0.7	9.0	0.5	0.3	0.3	0.5	Ξ	1.9	1.2	29.5	38.6	25.1	32.6	1.6	8.0	0.7	3.0	5.5	0.3	10.418
215977_x_at	X68285.1		glycerol kinase	1.9	6.0	.0.8	0.7	1.0	0.3	9.0	6.0	1.0	15.8	19.8	9.7	10.9	0.1	0.1	0.2	0.2	0.5	0.3	10.203
215783_s_at	X14174.1		liver-type: alkaline phosphatase	0.5	1.2	0.7	6.0	0.4	0.7	0.5	0	9.0	21.2	45.8	1.6	12.8	1.2	0.7	0.5	0.5	0.5	6.	10.167
217167_x_at	AJ252550		GK gene for glycerol kinase, exon 1	2,5	0.1	0.5	0.3	0.7	0.2	0.1	9.0	0.3	0.8	13.6	7.5	7.2	7.0	7.0	0.1	0.9	0.5	0.5	10.067
213349 at	A1934469		KIAA0779	0.0	0.7	0.9	1.4	1.0	1.9	1.4	2.3	2.1	15.6	15.3	19.5	23.8	2.2	7.4	7.5	7.0	7.2	7.7	9.7704

Mail						ပ္ဆိုင္ခ				F 62				70			No.	V aN							
Mail Control	٦	robe set	Accession #		Transcripts	plood		3 -	- 1	(smali)	Eo 1	~	Ī	(small)	Ne 1	Ne 2	(small)	(small)	<u>-</u>	CD4	800	CD14	6019	£	Ne S.I.
MARIONISTA CPPR polity gardide neceptor 2 0.1 0.2 0.4 0.7 0.7 0.9 0.7 0.8 0.7 0.8 0.7 0.8 0.7 0.8 0.7 0.8 0.7 0.8 0.7 0.8 0.7 0.8 0.7	١.٠	10789_x_at	L00692.1		carcinoembryonic antigen (CGM1)	1.1	0.4	1.9	1.4	1.5	2.0	1.5	2.5	0.4	19.4	18.6	22.6	15.5	0.7	1.6	6.0	5.0	0.3	1.1	9.2749
NM_ODSSSS PROFIGE PROFIGE NA 0.7 0.5 1.6 0.5 0.7 0.5 1.6 0.7 0.5 1.1 0.9 0.7 0.5 1.1 0.9 0.7 0.5 0.7 0.5 0.7 0.7 0.7 0.7 0.7 0.7 0.7 0.7 0.7 0.8 0.7	٠,٧	10772_at	M88107.1	GPR		0.1	0.2	9 .0	0.7	0.7	0.7	0.3	0.8	0.7	51,4	7.79	44.8	46.4	0.5	0.5	0.5	5.6	0.1	0.1	9.2661
NM_00195.1 R Fragman R Info (CDL6) 0.7 0.2 1.2 0.7 0.6 0.7 0.8 0.7 0.7 </td <td></td> <td>18978 s. at</td> <td>NM_018586.1</td> <td></td> <td>PRO1584</td> <td>0.7</td> <td>0.5</td> <td>9.1</td> <td>0.2</td> <td>0.5</td> <td>7</td> <td>6.0</td> <td>0.7</td> <td>0.5</td> <td>26.3</td> <td>20.5</td> <td>9.9</td> <td>8.8</td> <td>0.2</td> <td>0.7</td> <td>0.4</td> <td>1.1</td> <td>0.5</td> <td>0.4</td> <td>9.1536</td>		18978 s. at	NM_018586.1		PRO1584	0.7	0.5	9.1	0.2	0.5	7	6.0	0.7	0.5	26.3	20.5	9.9	8.8	0.2	0.7	0.4	1.1	0.5	0.4	9.1536
NW_001995.1 (https://doi.org/curier. Alignes. NW_001995.2 (https://doi.org/curier.org/curier. Alignes. NW_001995.2 (https://doi.org/curier.org/curier. Alignes. NW_001996.2 (https://doi.org/curier.org/curier. Alignes. NW_001996.2 (https://doi.org/curier.org/curier. Alignes. NW_001996.2 (https://doi.org/curier.org/curier.org/curier. Alignes. NW_001996.2 (https://doi.org/curier.org/curier.org/curier. NW_001996.3 (https://doi.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.org/curier.org/curier.org/curier.org/curier. NW_001999.3 (https://doi.org/curier.o	. •	04006_s_at	NM_000570.1	œ	Fc gamma R IIIb (CD16)	0.7	0.2	1.2	0.7	9.0	0.7	9.0	0.0	0.7	230.7	278.8	47.6	45.9	4.2	6.1	6.9	12.0	3.9	0.1	9.0769
NN_005062 male garm cell-associated kinase No. 1.3 6.7 6.8 6.7 6.8 2.8 2.8 4.7 3.2 2.4 4.8 48.8307.4 (CM)2 Invacious linear receptor binding belace. KGN/2 Invacious linear lineary belaces with the second lineary lineary belaces at the second lineary li	(4	07275_s_at	NM_001995.1		fatty-acid-Coenzyme A ligase. long-chain 1 (FACL1)	4.3	5.6	r.	11.5	19.4	5.3	5.7	5.9	3.7	93.8	127.8	92.8	69.1	0.7	1.0	1.1	8.3	6.0	1.7	8.9855
Augasco I (CNG investigle binding bedro-Z (L) 0.6 3.2 4.4 4.4 2.4 4.7 5.1 3.2 Augasco I (CNG investigle binding bedro-Z (L) 0.6 2.5 4.4 4.4 2.4 4.7 5.1 3.2 Augasco I (CNG investigle binding bedro-Z (L) 0.6 0.7 0.7 0.8 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.0 0.8 2.2 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1	"	20302 at	NM 005906.2		male germ cell-associated kinase	0.1	0.0	1.3	0.7	0.8	0.7	0.2	2.0	1.4	7.9	8.4	7.9	6.8	0.3	0.1	0.0	0.1	0.1	0.0	8.866
AF153820.1 ICN ANALOTE MANALOTE COMPAND DIVERSION AND ANALOTE AT MANALOTE COMPAND MANALOTE AT MANALOTE COMPAND MANALOTE COMPAND MANALOTE AT MANALOTE COMPAND MA		21803_s_at	AA883074		nuclear receptor binding factor-2	Ξ	0.8	3.2	4.0	5.9	5.8	2.8	4.7	3.2	24.7	25.2	33.2	29.3	0.1	1.5	5.	1.7	5.0	5.6	8.3788
A893469 (CAA0778) (AAA0778 (AAA0778) (AAA0789)		06765_at	AF153820.1	Ď.	channel Kir2.1	1.0	9.0	2.5	4.4	4.4	2.4	4.7	6.1	8.0	25.8	36.2	52.4	48.3	9.0	0.2	0.5	1.5	0.7	1.1	8.108
ALŞASFGO (ubiquitive) quantities and pydroxylase of the control of	. •	13351_s_at	A1934469		KIAA0779	0.0	0.1	6.0	2.0	1.6	9.0	9.0	2.3	2.7	7.9	5.9	20.1	18.0	0.5	0.5	0.2	0.1	0.2	1.2	7.9705
NW_O04688.1 CivP4F3) CivP4F3	••	14590_s_at	AL545760		ubiquitin-conjugating enzyme E2D 1	0.1	1.5	0.3	0.4	0.5	₹.	1.0	0.5	9.4	7.9	10.7	10.1	4.4	0.1	4.0	0.3	0.	0.4	6.0	7.8712
UB09391 R F cgamma receptor IIc2 1.9 1.0 1.7 2.1 3.3 8.1 16.1 2.8 4.2 NM.0108391 VML OB0167.1 Sylvaprotein regulator 2 2.1 1.5 1.0 1.5 1.6 1.5 1.6 1.8 1.1 1.4 1.6 1.4 1.6 1.4 VML OB0167.1 Sylvaprotein regulator 2 2.1 1.5 1.0 1.5 1.6 1.5 1.6 1.1 1.4 1.6 1.4 1.6 1.4 VML OB0167.1 Sylvaprotein regulator 2 2.1 1.5 1.0 1.5 1.6 1.5 1.6 1.1 1.4 1.6 1.4 1.4 1.6 1.4 1.6 1.4 1.6 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.6 1.4 1.4 1.4 1.6 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4 1.4	. •	06522_at	NM_004668.1		(CYP4F3)	0.0	0.0	9.0	4.8	1.0	3.0	5.4	11.7	9.4	46.0	51.5	56.9	43.0	0.1	0.0	0.0	9.0	9.0	0.1	7.5773
NM_018399.1 VNN3 protein NM_018399.1 VNN3 protein NM_000457.1 early development regulator 2	. •	10992_x_at	U90939.1	œ	Fc gamma receptor Ilc2	1.9	1.0	1.7	2.1	3.3	8.1	16.1	8.2	4.2	70.9	92.6	42.8	30.5	0.3	0.5	0.1	7.2	2.3	0.2	7.5239
NM_000427.1 spiry-development regulator 2 5.6 4.4 4.8 6.5 9.6 7.6 6.8 5.3 7.3 11. NM_000427.1 spirore kinase and beliable evident HSP70B	. •	20528_at	NM_018399.1		VNN3 protein	0.1	0.2	1.3	6.1	5.0	0.5	0.1	0.5	9.0	26.8	22.1	20.0	36.8	9.0	0.5	0.1	3.0	0.1	0.1	7.5216
NM_000167.1 gyvcarol kinase NM_10243.2 juluaninyl-peptide cyclotransferas NM_01963.1 myeloid cells 1	•	00919_at	NM_004427.1			5.6	4.4	8.8	6.5	9.6	9.7	6.8	5.3	7.3	109.5	89.0	52.9	61.4	2.5	6.9	7.0	10.0	3.9	6.5	7.5056
X51757 heat-shock protein HSP0B 1.1 0.2 0.8 0.6 0.9 5.9 8.3 2.1 4.1 0.0 0.4 0.0 0.4 0.7 0.9 0.7 0.0 0.4 0.3 0.4 0.0 0.4 0.7 0.9 0.7 0.4 0.3 0.4 0.2 0.5 2.9 0.7 1.4 1.5 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.4 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1<	. •	07387_s_at	NM_000167.1		glycerol kinase	2.1	.1.5	1.0	1.5	1.6	7	1.4	1.6	1.4	14.3	20.3	10.7	10.3	0.9	9.0	0.3	1.7	9.0	9.0	7.4603
NM_012413.2 gitularmive-peptide cyclotransferas 0.4 0.0 0.4 0.7 0.9 0.7 0.0 0.4 0.3 3.3 NM_018643.1 ritiggering receptor expressed on mayorid cells 1 ritiggering receptor expressed on mayorid cells 1 1.3 0.3 0.4 0.2 0.5 2.9 0.7 1.4 1.5 NA 0.9 0.7 0.0 0.4 0.5 1.4 1.5 NA 0.0 1.4 1.5 1.4 1.5 NA 0.0 0.0 0.1 0.1 0.1 0.4 0.4 0.2 0.5 0.7 0.1 <t< td=""><td></td><td>17_at</td><td>X51757</td><td></td><td>heat-shock protein HSP70B</td><td>7</td><td>0.2</td><td>9.0</td><td>9.0</td><td>6.0</td><td>5.9</td><td>8.3</td><td>2.1</td><td>1.4</td><td>46.3</td><td>41.2</td><td>25.1</td><td>25.8</td><td>0.4</td><td>6.0</td><td>9.0</td><td>3.7</td><td>9.</td><td>0.5</td><td>7.4139</td></t<>		17_at	X51757		heat-shock protein HSP70B	7	0.2	9.0	9.0	6.0	5.9	8.3	2.1	1.4	46.3	41.2	25.1	25.8	0.4	6.0	9.0	3.7	9.	0.5	7.4139
NM_018643.1 Imagenary decentant repressed on myeloid cells 1 Imagenary decentant repressed on myeloid cells 1 1.7 0.1 0.4 0.2 0.5 2.9 0.7 1.4 1.5 NA_000361.1 NM_000361.1 thrombomodulin or myeloid cells 1 thrombomodulin or myeloid cells 1 1.7 0.1 0.4 0.4 0.2 1.0 1.1 0.0 AM_000607.1 FLL20273 crossomucoid 1 (ORM1) 0.5 0.4 0.1 1.2 1.7 0.8 1.1 0.6 1.1 0.0 AM_019027.1 FLL20273 Cocal adhesion kinase po125(FAK) 2.9 1.0 0.9 1.5 0.8 0.9 0.4 1.7 0.8 0.6 1.7 0.8 0.6 1.7 0.8 0.6 1.7 0.8 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 0.8 1.7 1.7 0.8 1.7 1.7 0.8 1.7 <	. •	05174_s_at	NM_012413.2		glutaminyl-peptide cyclotransferas	0.4	0.0	0.4	0.7	0.9	0.7	0.0	0.4	0.3	30.1	47.7	36.4	24.7	0.7	9.0	0.0	4.6	0.1	£.	7.3246
NM_000561.1 thrombomodulin orusomucoid I (ORM1) 1.7 0.1 0.4 0.4 0.2 1.0 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1 1.1 0.1 0.2 1.1 0.1	. 4	19434_at	NM_018643.1		<u>5</u>	1.3	0.3	0.4	0.2	0.5	2.9	7.0	1.4	1.5	92.8	84.0	81.2	98.6	3.4	9.0	0.1	12.3	6.0	0.3	7.2368
NM_000607 1 corsomucoid I (ORM1) 0.5 0.1 0.1 0.4 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.0 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1		03887_s_at	NM_000361.1		thrombomodulin	1.7	0.1	0.4	0.4	0.4	0.5	1.0	1.4	1.1	16.8	13.0	13.4	1.3	0.7	9.0	0.4	1.1	9.0	0.5	7.1328
ANM_019027.1 FLJ20273 NM_019027.1 GTPase regulator associated with the focal adhesion vinase pol/25/FAK) NM_019027.1 GTPase regulator associated with the focal adhesion vinase pol/25/FAK) NM_019027.1 GTPase regulator associated with the focal adhesion vinase pol/25/FAK) NM_019027.1 GTPase regulator associated with the focal adhesion vinase paulogene. NM_019027.1 GTPase regulator associated with the focal adhesion vinase pol/25/FAK) NM_019027.1 FLJ2027.2 FLJ202/FAK) NM_019027.1 FLJ2027.2 FLJ202/FAK) NM_019027.1 FLJ2027.2 FLJ202/FAK) NM_019027.1 FLJ2027.2 FLJ2027	. •	05040_at	NM_000607.1		orosomucoid 1 (ORM1)	0.5	0.1	0.1	9.0	0.1	0.1	0.0	1.1	0.0	3.4	4 .4	4.3	0.7	0.1	0.3	0.1	0.4	0.0	0.5	7.0823
NM_019027.1 FLJ20273 NM_019027.1 GTPase regulator associated with the focal adhesion kinase pol/25/FKK) BEGT1084 (GTPase regulator associated with the focal adhesion kinase pol/25/FKK) NM_003064.1 finibitor (antileucoproteinase) (SLPI) X78713 carcinoembryonic antigen X78713 carcinoembryonic antigen NM_024850.1 FLJ21458 NM_0024850.1 FLJ21458 NM_00578.1 transducialise enhancer of spil 3 AIS67426 NM_000760.1 R receptor superfamily member NM_018340.1 FLJ311551 NM_018340.1 RLJ311551 NM_018340	. •	14681_at	A1830490		glycerol kinaso	0.5	0.4	0.1	1.2	1.7	0.8	1.1	9.0	1.2	7.8	14.2	24.0	12.1	6.0	0.0	0.1	1.9	1.0	0.3	9969
GTPass regulation states with the focal adhesion kinase pot/25(FAK) 1.2 1.4 3.5 5.9 5.1 6.5 5.1 8.5 7.1 4.5 6.5 6.5 6.5 7.1 4.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.1 7.2 7.2 8.5 7.2	• •	18035_s_at	NM_019027.1		FLJ20273	2.9	1.0	6.0	1.5	9.0	6.0	9.0	1.3	1.1	47.2	54.4	91.9	94.0	1.7	0.7	0.4	6.6	1.5	0.3	6.9074
NM_003064.1 Inhibitor (anillativoproteinase) (SLPI) 1.7 0.8 4.6 1.5 1.1 1.2 2.2 8.5 0.8 1.9 A/8713 Actionosome Arighmen	• •	05068_s_at	BE671084		GTPase regulator associated with the focal adhesion kinase pp125(FAK)	1.2	1.4	3.5	5.9	5.1	6.5	5.1	8.5	7.1	43.7	40.6	45.2	56.0	0.7	1.3	5.9	5.1	9.0	1.2	6.8934
State	. 4	03021 at	NM_003064.1		secretory leuxocyte protease inhibitor (antileukoproteinase) (SLPI)	1.7	0.8	4.6	1.5	1.1	1.2	2.2	8.5	0.8	16.2	19.8	13.9	8.7	1.8	9.0	1.2	1.7	1.1	2.1	6.7497
Control of the cont		16316_x_at	X78713		glycerol kinase pseudogene, chromosome 1	2.0	6.0	0.4	9.0	1.0	0.7	9.0	0.5	1.0	10.3	18.6	. 8.9	5.8	0.5	0.2	0.2	0.8	0.3	0.1	6.6675
NM_024850.1 FLJ21458 0.3 0.6 0.1 0.1 0.1 0.3 0.6 0.7 0.7 0.7 0.8 0.8 1.3 0.9 0.5 8.5 18.7 10.1 11 10.1 11 10.1 11 10.1 11 10.2 2.4 2.0 10.2 2.3 10.6 2.4 2.0 10.1 10.1 10.1 10.2 10.2 10.2 2.4 2.0 10.2 2.4 2.0 10.2 2.4 2.0 10.2 2.4 10.2 2.4 10.2 2.4 10.2		17209 at	X16454		carcinoembryonic antigen subdomains A and B	9.0	9.0	0.2	0.2	0.4	0.1	0.3	0.7	0.5	2.8	3.8	3.3	2.7	0.3	0.4	0.0	0.1	0.1	0.5	6.5382
NM_O15714.1 putative lymphocyte GOG1 switch 0.8 0.8 1.3 0.9 0.5 8.5 18.7 10.1 10.2	.,	20421_at	NM_024850.1		FLJ21458	0.3	9.0	0.1	0.1	0.1	0.3	0.8	9.0	0.3	8.9	7.2	9.1	7.0	1.2	9.0	0.0	0.1	0.1	0.2	6.4857
Al567426 KIAA1547 transducin-like enhancer of split 3 2.2 2.6 1.3 0.7 0.5 1.6 0.5 1.0 2.2 1.9 Al567426 KIAA1547 KIAA0329 0.7 0.0 0.1 2.6 2.1 1.8 1.5 2.4 2.0 granulocyte colony-stimulating factor immunoglobulin superfamily member NM_0018340.1 R_LLJ11151		13524 s at	NM_015714.1		putative lymphocyte G0G1 switch gene (G0S2)	0.8	0.8	1.3	0.9	0.5	8.0	8.5	18.7	10.1	118.0	. 25.0	39.6	92.5	1.7	0.8	0.1	6.0	0.2	3.4	6.4779
AB002295.1 K/AA0329 AB002295.		06472 s at	NM 005078.1			2.2	2.6	1.3	0.7	0.5	1.6	9.0	0.5	1.0	22.6	25.0	11.3	8.3	1.3	1.2	1.5	1.8	1.5	0.0	6.4334
AB002295.1 K/IAA0329 NM_000760.1 R receptor colony-stimulating factor MM_000760.1 R receptor with MM_000760.1 R receptor mannoglobulin superfamily member		12769_at	AI567426			1.4	1.5	0.5	1.3	2.3	Ξ	1.0	2.2	4.9	17.9	15.9	15.1	12.8	2.4	0.8	7.	1.3	1.2	0.5	6.4271
MM_000760.1 R receptor colony-stimulating factor of the colony-sti	. •	04307_at	AB002295.1		KIAA0329	0.7	0.0	0.1	5.6	2.1	1.8	1.5	2.4	2.0	10.3	11.2	15.4	12.8	1.9	0.1	0.5	0.2	0.1	0.5	6.3478
AF181660.1 WM78 NM_018340.1 FLJ11151 NM_021842.1 R Fcgarma receptor IIa (CD32)		03591_s_at	NM_000760.1	œ	granulocyte colony-stimulating factor receptor	0.1	0.0	1.0	0.1	0.2	2.3	9.0	2.4	1.0	220.3	187.2	134.9	112.1	0.2	8.0	0.2	25.5	0.1	0.5	6.2051
NM_018340.1 FLJ11151 1.5 1.2 1.2 1.2 1.8 3.3 3.2 2.6 1.8 NM_021642.1 R Fc gamma receptor IIa (CD32) 4.7 1.2 0.9 1.2 2.4 19.5 22.8 19.4 21.4 1 x1513.1 currorida dismutase 2 1.3 1.1 1.2 1.8 2.2 1.4 1.3 3.2 2.4	•	10210 at	AF181660.1		immunoglobulin superfamily member WM78	0.9	1.1	9.1	. 60	6.	7	6.1	1.7	2.0	1.1	7.0	13.7	14.1	4.	1.3	6:0	1.6	7	9.6	6.172
NM_021642.1 R Fc gamma receptor IIa (CD32) 4.7 1.2 0.9 1.2 2.4 19.5 22.8 19.4 21.4 1 x1513.1 superovide dismulase 2 1.3 1.1 1.2 1.8 2.2 1.4 1.3 3.2 2.4	. •	18610_s_at	NM_018340.1		FLJ11151	1.5	1.2	1.2	1.2	1.8	3.3	3.2	5.6	1.8	25.2	23.3	17.9	21.8	0.5	9.0	0.4	3.6	0.4	9.0	6.112
X151321 curporavide dismutase 2 13 11 12 18 22 14 13 3.2 2.4	- 1	03561_at	NM 021642.1	α	Fc gamma receptor IIa (CD32)	4.7	1.2	6.0	1.2	2.4	19.5	22.8	19.4	21.4	104.3	129.0	153.0	123.9	4.0	0.7	0.7	16.3	2.5	0.2	6.1015
A 13 132.1	••	216841_s_at	X15132.1	i	superoxide dismutase 2	1.3	1.1	1.2	1.8	2.2	1.4	1.3	3.2	2.4	20.5	44.6	35.8	42.2	5.6	8.0	0.7	2.0	0.7	1.1	6.0724

C. Neutrophil (Nc)-selective transcripts (2/7).

Figure 6F

Probe set Aα 207624_s_at NM 209850_s_at BC 20395_s_at NM 215965_x_at NM 20655_s_at NM	Accession #			ν C																		
# # # #	receion #			. p.	MC.	Ba	Ba 2 B	Ba 3		Eo 3	3 Eo 4			Ne 3	Ne 4							1
# # # #	# 1010000			ı	١	-	- 1	- 1	E0 1	Eo 2 (sm	- 1	lall) . Ne 1	Š	٠,	- 1	<u>ا</u>	5	80 5	CD14	CD19	٩	Ne S.I.
# # # # # # # # # # # # # # # # # # #	NM 000328 1	٠	retinitis pigmentosa GTPase regulator (RPGR)	0.0	9.0	1.1	2.0	1.2	7		2.1	2.8	7.1	.0	9		1.2	2 0.4	0.9	0.7	0.3	6.0348
<i>57 37</i>	BC005406.1		Cdc42 effector protein 2	0.2	0.5	1.3	0.5	1.0	0.2	0.4	0.5	0.4 1;	13.2	9	_	8.7 0.2	2 0.2		9.0	0.4	1.6	5.9313
5	NM 004994.1		matrix metalloproteinase 9	47.7	9.0	9.1	1.9	4.1	1.2	_	5.5	1.5	36.5 35.1	.1 36.8	8 21.9	.1	7 2.0	1,4	1.9	1.3	1.4	5.9311
	AA292874		glycerol kinase	1.2	0.5	9.0	0.1	1.2	0.2		1.2	1.2				7.4 0.2	2 0.1	1 0.2	1.3	0.5	0.0	5.9077
	NM_005668.1		sialytransferase 8	3.1	1.2	2.1	2.5	2.0	1.4	3.1	1.6	1.6 2	21.7 33.3	.3 6.8	8. 11.4	4 0.4	£ 0.9	9.1	2.6	7.	0.2	5.8688
₩	BC005980.1		ubiquitin-conjugating enzyme E2D 1	8 .	4.4	Ţ:	5.9	2.4	7.5	5.7	3.9	2.2 2.	29.3 31.2	.2 52.6	.6 25.4	4 0.2	2 . 2.0	1.4	5.7	1.3	2.3	5.8143
	NM 006224.1		phosphotidylinositol transfer protein (PITPN)	3.0	3.1	ę: 6	5.0	4.6	3.9	5.4	6.6	5.4 2	27.9 26.9	.9 32.9	.9 35.2	2 2.0	3.5	5 3.7	5.0	3.5	2.8	5.811
	U16120.1		placental taurine transporter	1.2	0.7	0.5	0.2	0.4	0.3	1.6	0.5	0.9	14.8 17.2	.2 2.8		2.6 0.0	0.1	1 0.3	Ξ	0.3	0.4	5.7909
209137_s_at BC	BC000263.1		ubiquintin c-terminal hydrolase related polypeptide	3.6	1.4	2.0	5.6	1.9	1.5	6.0	2.7	0.9	25.0 24.5	.5 37.6	.6 29.7	.7 3.5	5 5.0	3.5	3.5	2.7	3.0	5.7864
	NM_001815.1		carcinoembryonic antigen-related cell adhesion molecule 3 (CEACAM3)	1.7	4.4	2.9	2.7	1.9	3.0	2.6	2.8	2.1 14	14.3 16.5	.5 20.0	.0 13.7	.7 2.5	5 1.3	3 2.3	2.8	2.5	1.6	5.7572
	NM_000876.1	œ	insulin-like growth factor 2 receptor	6.4	2.8	0.2	0.8	1.8	1.4	5.4	6.8	5.4 8.	82.4 106.9	.9 80.1	1.77	(1)	7 2.9	9.87	8.4	4.8	14.8	5.7104
	NM_018169.1		FLJ10652	1.2	0.1	6.9	13.7	12.5	3.2	3.3		5.8 5	52.7 54.4	.4 82.8	.8 83.3	.3 2.9	9 8.7	7 11.8	3.2	10.8	1.3	5.6442
ā	NM_003003.1		SEC14 (S. cerevisiae)-like 1	5.7	2.5	12.8	28.8	22.6	50.6		•	29.5 13(-	-	-	.9 19.2			1.5	7.7	11.8	5.6423
32069_at AB	AB014515		KIAA0615	2.8	1.5	2.1	5.6	5.51 5.51	4.6	3.7	5.9	6.9	23.9 20.9	.9 35.1	.1 40.5	.5 1.0	3.4	3.3	2.6	2.1	1.5	5.6348
205896_at NM	NM_003059.1		solute carrier family 22, member 4 (SLC22A4)	0.8	1.2	1.0	0.1	1.	1.9	2.4	3.5	2.7	8.5 13	13.3 30.0		12.6 0.5	5 0.7	7 0.8	2.6	0.5	1.4	5.4612
	NM_015364.1		MD-2 protein	7.4	1.1	2.0	3.9	1.8	1.0	0.4		0.5 4	43.9 62.3	3 87.8	.8 56.0	.0 1.3	3.5	5 1.0	11.1	3.5	9.4	5.4459
ā	NM_003003.1		SEC14 (S. cerevisiae)-like 1	1.0	0.3	3.6	2.5	2.2	15.5	5.4 1;		8.4 41	40.2 57	57.3 54.9	6.09 6.	.9 3.8	9.0	8 1.3	2.7	3.3	2.4	5.4422
	NM_014844.1		KIAA0329	5.9	1.1	2.2	8.2	8.7	2.1			4.3	16.5 18.7	19.9	_	.0 3.4		1.7	1.8	1.3	2.5	5.4369
	NM_004347.1		caspase 5	0.3	1.1	0.1	0.0	0.1	0.1	0.3	0.2	0.5	4.2 6	6.2 4.6		2.9 0.3	3 0.3	3 0.4	0.8	0.1	0.3	5.379
203435_s_at NM	NM_007287.1		cD10, memprane metallo-endopeptidase	0.3	0.5	9.0	0.3	0.2	0.1	0.3	0.3	0.2 10	16.4 44	44.2 54.6	.6 42.8	.8 0.2	2 0.3	3 0.2	0.5	0.5	6.9	5.2591
	NM_006576.1		advillin	0.4	0.5	2.0	4.	8.0	1.0	<u>.</u>	1.0	-	8.4 6	6.9	.01 0.1	0.1 1.0	7 1.3	3 0.4	1.5	0.7	7	5.2541
	NM_014664.1		KIAA0615	1.9	1.2	1.3.	3.2	2.3	3.0	2.7	3.7	3.0 13	12.9 12	12.8 20.2		20.0 0.1	1 2.2	2 2.6	1.3	1.9	1.0	5.2381
201963_at NM	NM_021122.2		fatty-acid-Coenzyme A ligase, long-chain 1 (FACL1) CYCP1 interleukin 8 recentor	5.9	2.1	9.0	19.9	33.4	9.9	3.8 10	10.3	6.1	58.8 86	86.6 92.2	.2 80.2	.2 0.1	1.1	-	10.8	1.5	2.4	5.2374
207064_s_at NM	NM_009590.1	GPR	alpha	0.7	0.1	6.0	6.0	8.0	8.0	0.8	6.0	1.2	4.3 4	4.7 4.0		6.6 0.8	9 0.5	5 0.8	0.5	0.8	0.3	5.1905
220005_at NM	NM_023914.1	GPR	P2YX purinergic receptor GPR86 for UDP-glucose	9.0	0.0	0.1	0.5	0.1	19.7	9.6	25.1 1.	17.3 7	76.5 77.1	7.1 100.7	.7 98.6	6.1.9	9 0.1	1 0.2	12.3	9.0	0.0	5.1686
-	U90940.1	œ	Fc gamma receptor IIc3	4.6	0.3	7.3	7.6	15.1	7.6	19.5		6.6 11	-				1.4	9.0	14.0	7.2	1.0	5.1401
_	NM_007282.1		ring finger protein 13 (RNF13)	4.4	2.8	3.5	9.5	9.9	8.8	6.2	6.8	7.1 2	27.1 36	36.6 52.5	.5 35.7	7. 0.8	4.1	3.7	7.2	7.2	4.6	5.1305
209864_at AB	AB045118.1		GSK-3 binding protein FRAT2	5.8	1.3	4.4	9.5	8.0	13.4	16.9 2	27.3 27	22.6 9:22	93.2 97.2	.2 100.6	.6 104.2	2 1.3	3 1.9	9 2.9	10.2	2.5		5,1047
218319 at NM	NM 020651.2		pellino (Urosopnila) nomolog 1 (PELI1),	3.4	2.9	5.9	17.5	33.5	17.7	11.7 13	12.6	5.1 6	64.9 88	88.3 73.0	.0 80.2	12 2.4	6.3	3 2.9	13.8	12.6	Ξ	5.0403
at	NM_004900.1		phorbolin	9.0	7	1.0	1.2	5.6	1.3	9.0	3.9		19.6 11.1			0.5 1.5	5 0.2	2 0.4	0.4	1.0	4.2	5.0277
	NM_030938.1		DKFZp5661133	11.7	6.2	4.6	18.5	12.5	5.0	3.3	8.2	4.9 115.	5.5 117.2	.2 125.3	.3 122.5	.5 2.6	5.3	3 4.2	24.1	4.4	8.3	4.9858
	BC004395.1		apolipoprotein L	0.7	8.2	9.0	2.1	1.6	2.0	0.4	1.8	1.1	8.0 3	3.9 12.7	_	5.2 0.2	2 0.9	9 0.4	1.4	1.1	0.4	4.9332
204748 at NM	NM 000963.1		COX2 prostaglandin-endoperoxide synthase 2	5.5	16.2	9.0	2.4	2.0	5.	23	3.4	3.4	35.8 44.1	1.1 44.6	.6 91.8	. 6	4 0.6	5 0.2	4.3	0.3	0.5	4.9206
at	L32185.1		integral membrane protein	2.0	0.5	1.4	1.6	1.3	1.3	2.2		2.0 8							14.7	1.0	1.0	4.8845
	NM_005242.2	GPR	PAR2 proteinase activated receptor-2	8.0	9.0	6.0	0.2	6.0	6.0				11.8 15				3 1.5			0.3	0.7	4.8608
	A1934469		KIAA0779	9.0	6.0	1.2	1.5	1.2	1.2	1.5	1.5							9 1.5	1.0	1.3	0.8	4.8373
201888_s_at U8	U81379.3	œ	interleukin-13 receptor	0.1	9.0	0.3	0.2	0.0	1,7			2.0	8.6	19.0 15.	5.0 13	13.9 0.4	0.4	1		1.3	<u>.</u>	4.746

C. Neutrophil (Ne)-selective transcripts (4/7).

Figure66

cord
1.4 7.2
1.9 2.1
2.7 2.
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Figure 6.H

C. Neutro	phil (Ne)-selec	C. Neutrophil (Ne)-selective transcripts (5/7).																				
Probe set	Accession #	Transcripts	MC, cord blood	MC.	Ba -	Ba 2 (small) (Ba 3 (small)	Eo 1	Eo 2 (8	Eo 3 Eo (small) (s	Eo 4 (small)	- -	Ne 2	Ne 3 N (small) (s	Ne 4 (small) p	<u>.</u>	CD4	CD8 CD14	14 CD19	19 Fb	S. e	S.I.
204074 2 24	ALL DOCUMENTS	diotara acibaid Cha diotara somul	[٤	l	ı	l	-	٦	١	l	10.5	1	٦	Ĺ	4				۱,	Ι.	148
210594 x at	AF239756.1	myelin protein zero-like 1	1.2	0.5	9.0	1.5	1.2	0.1	0.4	0.8	1.9	15.3	13.9	1.7	10.9	0.3	0.2	0.6	1.0	0.7 2.8		3.9778
		protein phosphatase 1F (PP2C		,	;	;	,	:	;	;	•	;			:						_ '	Ş
203063_at	-		0.5	0.4	4.4	5.5	3.0	7	9.0	<u>.</u>	0.01	33.1	7.77	9.87	8.1.8						· ·	5 1
201392_s_at	BG031974 R	_	7.	0.7	0.1	0.0	0.1	5.6	3.0	6.	1.5	51.1	59.1	12.8	15.0		7.	4.0	3.8	2.2 7.0	e 	209
221477_s_at	BF575213		2.9	1.4	3.4	4.3	9.0	4.2	4.2	4.2	4.2	8.95	63.6	36.1	46.6	7.9		•			e-	.9562
206756 at	NM 019886.1	carbohydrate (N-acety/glucosamine 6-O)sulfotransferase 7	0.2	0.1	0.1	0.0	0.1	0.1	0.1	0.1	0.1	5.0	3.9	3.5	10.5	0.1	1.3	1.0	1.0	0.2 0.5	· ·	9516
	4	myxovirus (influenza) resistance 2																				
204994_at	NM_002463.1	(MX2)	1.3	2.3	5.4	7.8	8.2	9.5	9.1	15.2	12.3	49.8	46.6	9.77	80.8			•	5.6	4.8	<u>د</u>	.9478
201943_s_at	NM_001304.2	carboxypeptidase D	0.9	6.0	0.5	2.1	3.6	1.7	7.6	7.2	4.6	16.4	21.3	18.7	15.2	0.7				0.4 3.	د	926
216252_x_at	Z70519.1 R	CD95, Fas, APC	0.3	8.0	4.3	. 2.5	-	4.8	4.7	8.	1.7	18.5	31.2	10.3	8.4	2.2	3.8	2.4	1.9	1.4 3.9	<u>~</u>	1691
		B cell RAG associated protein	•	;		;	,	;	,	;	6	:		į		,		•	;		_	100
203066_at	NM_014863.1	(BRAG)		9.	6.9	7.	0	10.6	70.7	13.	7.07	0.70	65.9	35.6	80.9	e	~ .	0.0		7 .	2 (,000
212479_s_at	AL050139.1	FLJ13910	1.7	9.	7.	5.5	9.7	2.4	5.5	3.5	1.7	9.0	-0	80	8.6		2.3	2.4	2.2	e: 0	<u>۳</u>	.8422
209571_at	U03644.1	recepin	0.7	0.5	9.	3.3	2.7	2.2	1.3	5.6	2.7	8.9	7.8	12.8	10.9	9.0	1.0	-	1.2	ئة. د .	3.8	.8367
10000	, , , , , , , ,	CASP8 and FADD-like apoptosis	4	٠	4	96	•	,	9	,	000	11	0 77	45.7	47.6	7.	7 7		6.7	40		28332
Z11317_5_8I	AF04 1451.1	C-type (rainim dependent)	9	0.	ņ		•	ò	0,	3.5	9		?	4.5	2							3000
		carbohydrate-recognition domain)																			_	
209732 at	BC005254.1	lectin, superfamily member 2	 1.0	5.9	10.0	18.6	23.6	9.0	10.0	8.9	7.3	72.9	77.0	99.7	75.5	1.1	14.4 21	21.2 10	0.2 16.	1,3		3.8003
1		clone 24450 RING zinc finger protein																				;
201779_s_at	AF070558.1	RZF	14.8	6.0	13.0	13.3	13.9	21.8	27.0	25.3	13.1	88.2	108.1	116.5	81.8			1.3	25.7	5.7 8.5		3.7993
212441_at	D86985.2	KIAA0232	5.4	5.9	3.5	8. 6.	10.3	0. 0.	6.9	1 .5	1.3	31.9	30.1	30.8	49.8	5.5	7.0	0.		_	_	3.7868
205920 at	NM 003043 1	Solute carner tamily b, member b (SLC6A6)	1.2	14	60	0.0	0.0	1.1	5.2	0.7	1.4	33.1	26.7	2.7	4.0	0.3	0.1	2.	2.6	0.1 0.		3.7836
1070007	1000010	(Second)	: ;		? ;					;												3 70 4 6
213596_at	AL050391.1	DKFZp586A181	0.7	0.7	2.3	3.6	3.0	1.2	7.7	-	1.2	10.6	6.01	6.21	50 50	7.7	· [2			c o
209508_x_at	AF005774.1	apopinais	1.7	2.7	10.8	9.7	11.5	4.9	6.3	6.0	8.4	37.0	34.8	34.7	40.8	2.7	4.0	_	6.3	4		3.7451
218115 at	NM 018154 1	FI.110604	6.0	2.2	2.4	1.5	6.0	1.8	2.1	1.3	1.1	10.5	1.6	6.7	7.7	1.1	1.2	2.2	2.1			435
215652 at	AK024382.1	FLJ14320	0.2	0.1	2.0	0.3	1.6	0.3	0.3	1.2	0.7	5.5	4.5	3.4	2.5	0.1	0.3	0				3.7249
212561 at	AA349595	RAB6 interacting protein 1	10.0	6.1	10.2	12.4	1.4	18.7	14.9	12.1	10.0	66.3	0.09	47.6	37.1			_	3.9	5.9 10.2		3.6962
100000	100000	23004417	96	ò	;	9	č	;	,	č	0	-	12.5	•	00	,	0	1,			_	3,6908
20a too_ar	NM_U14903.1	KIAAUSOS	0.0		3 ;	0.5	5 6	2 6	2 6		9 6		3 :	,	9 .						_	000
221874_at		_	S.	S.3	<u>`</u>	Ξ.	ر د د	9.	9.9) ·	6.0			- ·	7 .	ر د د		٠,				000
217207_s_at	AK025267.1 R	butyrophilin like receptor	1.2	1.1	4.4	2.1	1.7	1.2	1.9	9.0	2.4	10.9	3.4	7.2	- .				0.9	1.5		3.6795
212579_at	AA868754		1.3	1.6	5.3	8.6	15.5	7.3	7.8	11.3	8.3	32.8	42.9	40.0	39.7	5.0	5.4	5.7	-	0.5		3784
212657_s_at	AW083357	IL-1receptor antagonist IL-1Ra (IL-1RN)	48.8	2.3	6.0	0.5	9.0	2.7	6.0	1.5	1.3	35.2	37.0	48.2	35.5	9.0	0.8	0.3	9.3	0.4 0.	0.6 3.6	3.6759
202392 s at	NM 014338.1	phosphalidyserine decarboxylase	3.4	2.7	6.2	11.7	8.9	3.4	4.7	5.4	4.7	32.8	31.0	22.0	32.3	0.8		4.4	4.2	1.4		3.6753
100000	0 110000			,					0	,	,		;	•	4		,	_			3.6	2733
2001//_s_at	NM_000045.2	arginase (ARGT) LPS-induced TNF-alpha factor	6.0	3	7.7		0.0	8.	9	7	ò	n o	3	ë	9.0	3					,	7
200706_s_at	NM_004862.1	•	31.2	39.0	20.7	9.49	62.5	15.6	13.4	31.8	21.7	173.2	148.5	172.0	150.2	•			14.3	•-	18.2 3.6	9699
212478_at	AL050139.1	FLJ13910	0.1	0.1	0.5	2.3	3.5	1.4	8.0	1.5	1.0	3.7	3.0	4.8	6.5		0.3				۲,	9623
218660_at	NM_003494.1	dysferlin	2.5	0.5	4.5	0.5	0.5	1.7	0.2	5.6	1.1	48.5	51.8	44.5	20.4	3.3		1,7 10	9.01	0.2 1.	.,	.919
211982 x at	AL546600	exportin 6	9.6	6.7	14.9	31.8	27.8	24.6	24.0	34.0	24.0	107.2	101.1	9.98	90.2	•	-	17.4			۲,	3408
		guanyfate binding protein 2,	,	;	1	;	;	;	;	•		į	;		i	,				,	_	;
202748_at			2.4	5.3	3.7	9.0	9 ./	3.3	1.7	3.0	3.3	L.82	7.05	45.0	24.8	7.7			۵.		_	3.0111
220088_at	736.1	GPR C5a receptor	2.2	2.3	15.4	24.6	24.8	17.1	15.6	6 0	15.0	90.5	104.5	84.4	91.1	2.3	1.2	0.5 25	5.6		0.4 3.6	3.6036
202890_at	T62571	microtubule-associated protein 7	0.9	0.1	0.7	4.4	1.1	0.1	0.0	0.5	0.1	2.5	3.1	3.2	7.	0.0				-	-	3.6032
220987_s_at	NM_030952.1	DKFZP434J037	1.4	0.3	3.9	4.9	4.7	9.6	9.3	14.1	11.7	43.0	35.6	57.8	57.6	5.3	4.7	4.8 11	11.1	3.3 2	2.0 3.5	5748
																					l	

FigureGI

Probe set	Accession #	Transcripts	MC. cord blood	MC. tung	Ba 1	Ba 2 (small)	Ba 3 (small)	Eo 1	Eo 2 (Eo 3 E (small) (Eo 4 (small)	Ne 1	Ne 2	Ne 3 (small)	Ne 4 (small)	J.	CD4	CD8	CD14 C	CD19 F	a Z	Ne S.I.
207072 at	NM 003853.1	interleukin 18 receptor accessory protein (IL18RAP)	5.9	0.9	2.2	3.4	4.1	7.2	9.1	2.8	1.7	13.4	7.6	20.9	5.6	2.4	2.8	3.1	0.4	9.	0.3	3.5722
215719_x_at		CD95, Fas. APO-1	9,4	9.0	3.1	2.7	1.6	2.2	5.2	8.	1.7	19.0	31.9	4.1	9.4	1.0	4.5	2.2	9.6	:	4.4	3.5657
218404_at	NM_013322.1	sorting nexin 10	4.9	1.2	9.2	2.0	3.7	0.7	0.1	2.1	0.7	28.5	31.8	34.8	33.8	0.3	1.2	6.1	9.0	6.5	0.3	3.563
219394 at	NM 024419.1	phosphatidylglycerophosphate synthase (PGS1)	3.4	3.0	1.7	4.2	3.5	2.4	3.8	6.1	9.5	17.2	16.9	17.5	16.3	1.0	2.1	2.6	2.4	1.5	1.7	3.5544
216913_s_at		KIAA0690	0.1	0.3	0.1	0.1	0.1	1.9	2.5	2.3	2.8	8.9	8.3	7.5	10.9	0.1	0.1	0.2	2.5	0.1	0.0	3.5402
205118_at	M60626.1 GPR	R formytpeptide receptor 1	0.1	0.5	9.0	0.1	1.1	0.1	0.4	9.0	0.1	3.9	7.0	3.2	7.8	0.5	0.1	0.1	1.4	0.1		3.5295
210564_x_at	AF009619.1	FLAME-1-delta	1.4	1.5	5.3	9.9	7.0	4.3	4.2	5.9	3.5	21.2	30.3	18.4	19.6	2.1	5.9	4.0	4.6	3.0	_	3.5148
213607_x_at	BE551347	KIAA0134	1.3	1.3	5.9	2.0	1.7	8.7	15.8	3.5	4.3	42.7	42.5	1.1	15.3	9.0	1.3	1.9	6.2	1.7		3.514
203888_at	NM_000361.1	thrombomodulin	6.0	9.0	0.1	0.1	0.3	0.2	0.4	0.8	0.1	6.1	6.4	4.6	0.2	0.3	0.3	0.1	9.0	0.5	0.1	3.5126
210233_at	AF167343.1	receptor (P)	0.5	0.1	0.0	0.1	9.0	8.0	9.0	9.0	0.7	4.8	3.0	5.0	2.3	9.0	0.0	0.0	0.0	0.0	0.3	3.5023
204959_at	NM_002432.1	antigen.	1.2	9.0	24.2	20.0	26.2	17.5	17.9	49.8	22.3	249.8	290.2	186.0	217.0	3.4	1.6	9.0	9.99	2.5	0.0	3.4903
217967_s_at		niban	6.3	3.7	16.3	45.4	35.9	27.9	29.5	43.0	30.9	107.4	115.3	117.9	110.3	2.6	7.5	6.6	4.9	3.4		3.4847
221763_at	A1694023	топе	0.8	1.5	6.1	4.6	5.5	5.5	6.9	6.6	9.0	28.3	32.9	34.0	42.7	1.1	3.7	£.4	5.1	9.8	3.0	3.4777
207857 at	NM 006866.1	leukocyte receptor, subfamily A (with TM domain), member 2 (LILRA2)	0.4	0.5	3,6	8. 5.	5.3	4. 5.	6.9	9.	6.3	33.5	34.3	46.2	50.8	8.	0.0	0.1	11.6	0.5	0.1	3.4773
220740 s at	NM 005135.1	solute carrier family 12 member 6 (SLC12A6)	1.6	0.5	2.5	3.3	3.4	3.5	5.6	3.2	3.2	12.6	19.0	10.7	14.4	4.0	2.9	2.2	2.9	3.5		3.4722
217739_s_at		pre-B-cell colony-enhancing factor	7.1	1.7	6.6	46.0	61.0	18.9	13.9	12.9	9.3	112.4	120.5	8.06	93.4	1.2	2.3	4.4	17.6	2.1	3.1	3.4242
205041_s_at	1 NM_000607.1	orosomucoid 1 (ORM1)	0.3	1.0	0.4	0.1	0.1	0.0	0.1	1.0	0.1	5.6	3.6	2.7	0.4	0.1	0.5	0.1	0.5	0.0		3.4189
214784_x_at	I BE966299	exportin 6	7.4	9.0	10.0	23.7	23.9	13.0	16.7	23.3	20.5	60.3	8.79	57.2	8.09	5.9	8.7	7.8	9.9	5.9	5.4	3.4155
217985_s_at	1 AA102574	bromodomain adjacent to zinc tinger domain, 1A	1.5	1.4	8:	5.6	3.8	7.4	5.1	7.3	4.7	16.9	19.2	22.1	24.2	2.7	2.5	2.3	3.5	2.4	0.9	3.3999
212598_at	A1806395	KIAA0993	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	5.5	7.3	3.1	3.2	0.2	0.1	0.0	1.0	0.1	_	3.3933
219053_s_at	t NM_017966.1	FLJ20847	2.6	2.0	6.1	2.3	2.3	5.6	2.1	3.0	2.8	11.3	9.7	22.2	34.1	1.6	0.5	6.0	4.7	1.3	_	3.3686
217475_s_at	t AC002073	15N1	9.0	0.5	9.0	0.5	7 .0	1.3	£.	0.5	1.2	5.9	8.6	9.	1.4	0.7	0.3	0.1	0.5	7.0	0.2 3	3.3448
. 46323_at	AL 120741	reticulum nucleoside diphosphatase	4.4	3.2	4.2	8.3	7.2	5.3	. 6.7	11.4	6.7	20.0	21.5	33.9	26.3	2.5	8.2	3.6	4.3	3.4		3.3028
201965_s_at	1 NM_015046.1	KIAA0625	2.7	3.1	3.3	8.5	8.3	4.9	5.8	7.4	7.5	19.5	18.5	18.7	8.72	2.2	3.9	3.4	4.6	5.6	_	3.3019
203628_at	NM_000875.2 R	insulin-like growth factor 1 receptor	0.5	0.1	3.7	3.0	4.0	8.9	4.9	2.7	5.8	18.8	23.4	8.9	20.6	9.1	0.1	2.8	2.4	2.0	3.5	3.2989
202193_at	NM_005569.2	,	8.	1.5	6.1	5.6	9.6	6.4	4.8	11.3	8.9	17.0	20.5	36.5	28.2	0.1	1.3	1.	9.0	0.5	9.0	3.2887
203042 at	NM 002294.1	lysosomal-associated membrane protein 2 (LAMP2)	2.4	3.1	Ξ	1.7	1.3	5.9	6.4	12.4	10.7	21.3	27.0	1.4	35.4	9.0	9.0	0.5	3.2	9.0	9.4	3.282
220326_s_at	NM_018071.1	FLJ10357	5.7	4.5	1.9	1.2	1.9	6.3	8.9	5.2	8.5	25.3	28.1	19.2	21.0	6.0		0.7	6.4	0.1		3.2759
212470_at	AB011088.1	sperm associated antigen 9	4.0	5.9	3.3	6.3	7.2	4 .	7	4.7	6.9	13.2	17.0	18.1	24.0	1.2	2.5	2.2	3.9	3.0	5.4	3.2688
211133_x_at	AF009643.1	ipt 5	1.9	4.4	8.0	1.5	1.3	3.1	2.7	3.8	2.1	42.1	42.2	27.0	25.1	2.7	6.0	0.5	10.2	0.4		3.2526
219313_at	NM_017577.1	DKFZp434C0328	0.0	8.0	0.5	0.0	0.4	0.2	-6	1.0	0.0	3.8	3.7	3.1	5.6	0.2		0.1	0.0	1.2		3.2404
221149_at	NM_018485.1 GPR	R G protein-coupled receptor GPR77	0.2	0.7	0.8	0.7	1.2	0.7	6.0	6:0	9.0	3.3	3.2	4.2	5.9	0.0	0.3	0.2	4.0	0.1	0.3	3.2282
203433_at	NM_006441.1	synthetase	1.4	1.8	1.5	2.5	1.6	3.1	4.2	3.5	£.	15.8	1.4	9.6	9.0	1.6	1.8	1.9	5.9	1.7	1.5 3	3.2181
214486_x_at	t AF041459.1	FADD-like apoptosis regulator	6.1	2	10.7	5.0	6.5	4.2	5.3	4.3	9.0	26.3	16.2	25.3	24.2	2.4	3.7	20	5.3	5.3	┥	3.2114

C. Neutrophil (Ne)-selective transcripts (7/7).

Statistical composition of the property of t	Probe set	Accession #	Transcripts	MC. cord blood	MC. lung	Ba I	Ba 2 (small)	Ba 3 (small)	Eo 1	Eo 2 (Eo 3 E (small) (Eo 4 (small)	Ne 1	Ne 2	Ne 3 (small)	Ne 4 (small)	į	CD4	CDB	CD14 (CD19	æ	Ne S.I.
Note Control	209222_s_at		oxysterol binding protein-like 2	1.3	0.2	2.7	8.8	8.0	5.4	5.4	7.7	7.3	17.2	18.4	22.3	23.9		2.3	3.4	8.2	5.0	£.	3.1877
	202334_s_at		ubiquitin-conjugating enzyme E2B	3.3	3.4	8.8	15.5	12.1	5.3	7.2	14.7	12.8	20.9	25.6	43.6	38.1	4.0	4.5	3.9	2.5	4.9	4.6	3.1871
Activities Linking L	203266_s_at		mitogen-activated protein kinase kinase kinase 4	3.7	2.4	4.9	8.2	6.3	2.5	4.8	6.8	2.0	14.7	15.2	27.4	56.9	2.8	2.4	5.6	2.7	2.3	2.7	3.1804
Mail Control	58780_s_at		FLJ10357	5.5	5.4	1.5	3.9	3.0	7.8	9.6	12.8	16.4	28.0	27.6	46.7	44.1	1.2	9.0	0.1	6.5	0.5	2.7	3.1803
NALOMENIA TRANSPANCIANCE CONTINUE MICRORANI NECONARY C	210582_s_at		LiM domain kinase 2	3.8	1.9	4.3	5.1	8.4	8.7	10.3	10.8	9.4	33.9	36.5	30.8	23.8	1.3	1.7	æ.	1.9	6.0	1,4	3.1609
Na, Original Na, Ariginal Na, Ariginal Na, Ariginal Na, Original Na,	214766_s_at	AL080144.1	ELYS transcription factor-like protein TMBS62	1.3	0.5	5.0	3.1	4.2	1.9	4.4	2.1	2.4	9.4	9.5	6.6	11.3	3.2	1.3	9.0	0.5	1.0	1.3	3.1414
NALONGE	202266_at	NM_016614.1	TRAF and TNF receptor-associated protein (AD022)	69	6.2		23.7	20.9	9.6	6.6	16.5	14.0	36.1	56.5	64.5	57.5	4.5	7.7	9.4	5.3	8.2	6.3	3.1402
MAIOCRGRI Direct Once Proteins According MAIOCRGRI Direct Once Proteins MAIOCRGR Direct Once Proteins Direct Once Proteins Direct Once	203278_s_at		BRAF35/HDAC2 complex (80 kDa)	5.1	2.8	2.7	14.8	11.0	5.7	5.0	7.4	4.9	21.0	16.6	31.0	30.6	0.1	3.0	5.5	3.4	2.7	3.0	3.1382
MACOSTORIST DIGNET CONTINUED GROUNDS MACOSTORIST MACOSTORIST MACOSTORIST DIGNET CONTINUED GROUNDS MACOSTORIST MACOSTORIST MACOSTORIST DIGNET CONTINUED GROUNDS MACOSTORIST																							
MALODISSI Destrictive temply 11 12 12 13 13 13 13 13	207291_at	NM_024081.1	protein 4	0.1	1.2	1.3	1.4	1.1	6.0	0.2	0.4	0.2	3.8	6.7	4.5	8.5	0.5	0.8	0.1	1.9	1.1	9 .0	3.1336
MAYOSTEGS A STATE MAYOSTEGS A STATE MAYOSTEGS A STATE	213229_at	BF590131	Dicer1, Dcr-1 homolog (Drosophila)	8.5	5.5	19.5	12.9	15.4	15.9	9.7	19.1	16.0	35.2	39.0	4.19	69.4	6.2	6. 4	۲. ز دن	15.2	12.2	5.0	3.1306
Accordant Commentarion Comment	204204_at		solute carrier family 31	1.7	. :		6.3	8.0	. :	6.0			22.6	23.8	31.3	36.0	2.0	7.7	2.7		7.0	5 6	3.1262
A GORGEAL Impropried in successing 14	201364_s_at		clone 17.6 immunoglobulin-like	8.5	į	4 . 4	7.7	S:	e G	6.13	9	2	F	- 76	2.17	7.73	7.7	n	5.	7.7	Y	0	3.1233
MAYORSER	210784_x_at		transcript	1.4	0.1	2.1	1.2	2.4	2.5	2.2	2.8	2.1	52.1	48.8	31.8	36.3	8.0	0.5	0.5	13.3	0.3	0.5	3.1236
MAY	202625_at	AI356412	related oncogene homolog (LYN)	6.9	6.9	5.0	18.2	14.9	20.7	21.7	27.7	28.6	74.7	76.3	77.5	75.7	7.5	1.7	6.0	17.1	23.6	0.5	3,1131
Purple P	221895_at	AW469184	hypothetical protein MGC26706	2.2	0.5	5.4	5.2	5.8	2.5	3.4	7.6	5.1	14.8	12.0	24.6	19.2	5.0	2.4	2.1	5.2	1.5	9.1	3.11
Figure Proposed	37384_at	D13640	rocompanomity	2.5	3.0	3.4	89	3.8	6.7	9.9	7.6	9.9	25.2	17.9	21.3	29.4	5.6	2.5	2.5	7.4	4.8	8.8	3.0985
NN_OOD5451 R Tollide receptor 2 18 12 18 18 18 18 18	206608_s_at		interacting	1.1	4.	1.2	0.8	1.0	=	1.0	0.5	4.	6.1	3.4	5.3	4.7	1.5	6.0	9.0	£.	0.5	0.7	3.091
NM, 000043.1 P. CO96, Fee, APO.1 2.0 1.1 5.9 11.0 8.9 5.4 4.0 6.8 7.0 2.0 2.7 2.5 2.0 2.9 2.0 2.	204924_at	NM_003264.1	Toll-like receptor 2	1.8	0.1	2.1	8.2	9.7	2.8	0.4	9.0	1.4	52.9	88.0	94.4	8.66	1.6	1.3	0.1	26.3	6.0	0.4	3.0909
MAIOGRAFIA MAI	204781_s_at		CD95, Fas. APO-1	5.0	1.1	5.9	11.0	8.9	5.4	4.4	6.8	7.0	50.6	27.1	27.0	29.1	0.3	5.2	5.3	4.0	2.0	3.8	3.0827
NM_00015661 FLAME NM_00015661 NM_00015661 NM_00015661 NM_00015661 NM_00015661 NM_00015661 NM_00015661 NM_00015621 NM_0001661	212606_at	AI806395	KIAA0993	2.0	2.0	0.0	0.7	9.0	0.7	0.0	0.1	0.1	18.0	14.3	25.0	56.9	0.1	0.5	0.3	9.9	0.0	3.4	3.0797
NM_0004161 Interceloral architection A 2.5 3.2 6.4 4.8 10.7 9.7 18.5 12.1 29.3 34.1 44.7 46.1 0.5 1.2 1.7 8.3 0.2 2.9 1.3 NM_0005172 (ICAM3) Interceloral architection A 2.5 3.2 6.4 4.8 10.7 9.7 18.5 12.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 1.8 3.1 3.	211316_x_at	AF009616.1	FLAME-1	3.1	5.9	20.1	19.4	31.9	7.6	4.9	9.02	19.1	28.8	14.1	72.8	91.0	4 .	8.9	8.7	10.3	5.5	5.6	3.0752
NM_0051622 (CAM3) NM_0051622 NM_0051622 (CAM3) NM_0051622 NM_0051622 NM_0051622 NM_0051622 (CAM3) NM_005162 NM_0051622 NM_005162 NM_	203045_at	NM_004148.1	lar adhesion molecule	4.4	2.5	3.2	6.4	4 .8	10.7	9.7	18.5	12.1	29.3	34.1	44.7	46.1	0.5	1.2	1.7	86 E.	0.5	5.9	3.0746
NA_0007172 Carbonic anhydrase IV (CA4) CA2 CA6 CA CA CA CA CA CA C	, 204949_at	NM_002162.2		3.0	3.7	5.7	46.3	34.5	38.9	51.6	94.5	93.7	161.1	168.3	239.1	241.1	4.5	13.6	21.0	9.71	19.9	Ţ.	3.0617
ANY-30554 metalio phosphoesterases NM_002350.1 rejeted romogene formogene formogene viral metalio phosphoesterase NM_002350.1 rejeted romogene formogene viral metalio phosphoesterase NM_002350.1 rejeted romogene formogene viral metalio (SICS) superfamily NM_0052649.1 formation of the metalio phosphoesterase NM_005250.1 rejeted romogene formogene viral metalio (SICS) superfamily NM_005250.1 showing a viral metalio (SICS) NM_005550.1 showing forming for	206208_at	NM_000717.2	carbonic anhydrase IV (CA4)	0.5	9.0	0.1	0.1	0.1	1.1	0.7	1.0	9.0	5.6	5.2	2.8	2.4	1.2	0.1	0.1	0.1	0.1	0.1	3.0572
NM_0002350.1 immunoglobulin homolog (LVM) NM_0002350.1 immunoglobul	213727_x_at	AI743654	metallo phosphoesterase	2.1	1.7	5.0	15.0	12.1	6.8	13.7	25.4	23.9	49.9	28.3	59.9	75.8	0.7	4.5	9.9	5.7	5.4	3.7	3.0554
International control of the contr	202626_s_at	NM_002350.1	related oncogene homolog (LYN)	15.5	9.5	12.1	12.7	14.8	32.6	31.6	54.0	41.0	10.1	115.1	122.7	127.2	18.4	2.1	1.5	36.8	33.0	9.0	3.0541
Protein kinase C and casein kinase NM_007729.1 substrate in neurons 2 (PACSIN2) 2.0 1.0 1.3 3.7 3.6 20.1 15.3 19.5 33.9 35.3 61.3 72.5 76.1 94.4 24.7 1.9 1.0 0.2 1.1 1.3 1.0 0.5 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0 1.0	206420_at	NM_005849.1	immunoglobulin superfamily. member 6 (IGSF6)	-	0.3	0.2	0.4	0.7	6.	1.2	7	0.1	17.5	24.2	30.6	33.3	4.	9.4	0.4	8.4	0.3	0.3	3.0442
Additional problem of the protein 267 and the protein with a second and	201651_s_at	NM_007229.1	protein kinase C and casein kinase substrate in neurons 2 (PACSIN2)	21.4	17.4	4.6	24.8	20.1	13.3	19.5	33.9	35.3	61.3	72.5	1.91	94.4	24.7	4.9	4.4	8.6	6.1	8.0	3.0391
v.raf-Inturine Following layerial viral v.raf-Inturine Following layer v.raf-Inturine Following layer v.raf-Inturine Following layer v.raf-Inturine Following layer v.raf-Inturine	219540_al	AU150728		2.0	:+	1.3	3.7	3.6	2.0	1.5	4.4	3.7	6.7	6.9	9.5	6.6	1.0	2.1	£.	1.0	9.1	0.1	3.0361
AAB10268 kinase	201244_s_at			6.0	5.4	12.4	32.8	33.1	14.9	19.0	27.3	25.7	60.4	58.5	89.2	86.2	2.2	9.6	1.3	12.8	9.2	7.3	3.0329
AF007555.1 phosphatase NM_006526.1 interperior-induced protein with relations perior in character protein carbonic anhydrase IV (CA4) 0.7 1.0 0.8 0.6 0.6 0.6 0.6 0.6 0.6 0.7 1.1 1.5 0.7 1.0 0.8 0.6 0.6 0.7 1.1 1.5 0.7 1.0 0.8 0.6 0.6 0.6 0.7 1.1 1.3 1.4 1.7 1.3 1.4 1.7 1.3 1.4 1.7 1.3 1.4 1.7 1.4 1.7 1.4 1.7 1.3 1.4 1.7 1.7 1.7 1.7 1.7 1.7 1.7 1.7 1.7 1.7	203265 s at		proteín	7	1.2	8.8	6.4	4.5	1.7	5.6	3.5	3.1	12.8	12.8	10.4	16.8	1.4	4.	<u></u>	<u></u>	2.5	1.5	3.028
APOV535.1 prospirates NM_006526.1 interferon-induced protein with NM_006717.2 carbonic anhydrase IV (CA4) 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.	1000000		IAR receptor-like protein-tyrosine	. 6	6	•	;		3	3				ć	;	Ċ	č	;	6	2	2		3 0360
interferon-induced protein with 2.0 3.0 4.9 6.0 3.5 4.3 12.4 4.4 3.0 24.2 27.0 15.4 12.0 0.3 1.8 1.2 6.2 1.4 2.6 NM_000717.2 carbonic anhydrase IV (CA4) 0.7 1.0 0.8 0.6 0.6 2.2 2.1 1.5 2.1 13.8 7.4 4.7 3.5 2.1 1.0 0.5 0.9 1.0 0.5	203030_s_at		prospinalase zinc finger protein 217	0. 6.	2.6	3.0	18.4	9.4	7.6	9.0	14.7	14.9	37.3	45.1	24.4	27.0	1.2	4.6	5.3	. <u>.</u>	5.5	3.6	3.0158
NM 0007172 carbonic antydrase IV (CA4) 0.7 1.0 0.8 0.6 0.6 2.2 2.1 1.5 2.1 13.8 7.4 4.7 3.5 2.1 1.0 0.5 0.9 1.0 0.5	204747 at	NW 001549 1	protein	2.0	3.0	6.9	0.9	3.5	4.3	12.4	4.4	3.0	24.2	27.0	15.4	12.0	0.3	8.	1.2	6.2	4.	2.6	3.0143
	206209_s_at		carbonic anhydrase IV (CA4)	0.7	0.7	0.8	9.0	9.0	2.2	2.1	1.5	2.1	13.8	7.4	4.7	3.5	2.1	5	0.5	6.0	9	0.5	3.0112

D. Mast cell (MC)-selective transcripts (1/2).

					l																	
			MC.	Ş		200	200		-		-		e on	No.N	•							
Probe set	Accession #	Transcripts	plood	lung.	S -	_	(small)	Eo 1	Eo 2 (1 2	2 (small)		(smail) pl	ដ	CD4 CC	CD8 CD14	14 CD19	E.	MC S.I.	_:
217023_x_at	AF099143	tryptase beta	169.4	118.7	0.7	1.1	0.4	0.1	0.1	0.5	0.1	0.5	0.1	9.0	0.4 0	0.3 0	0.1 0	0.1	0.2 0.	1 0.2	202.148	48
215382_x_at	AF206666.1	tryptase beta	168.1	108.4	1.7	0.7	9.0	0.5	0.5	0.5	0.1	0.1	0.3	0.1	0.1 0	0.7.0	0.1 0	0.1 0	0.1 0.1		172.668	89
204041_at	NM_000898.1	monoamine oxidase B	23.5	46.9	0.5	0.7	0.1	0.1	0.5	9.0	0.3	0.3	0.1	0.2	0.1.0	0.2.0	0.2 0	0.1 6	0.2 0.1	1 0.2	136.311	=
210084_x_at	AF206665.1	tryptase alpha	131.1	92.3	0.7	1.2	1.2	0.5	0.1	0.1	0.1	0.1	0.1	0.1	0.1 0	0.6	0.1 0	0.0	0.1 0.1	1 0.1		92
216474_x_at	AF206667.1	tryptase beta	210.0	120.9	2.4	5.0	1.4	1.2	0.4	0.2	0.1	0.5	1.1	0.1	0.1	1.8	0.8 0	0.9	0.9 0.1	1 0.1	84.3375	75
205683_x_at	NM_003294.2	tryptase beta	195.5	95.3	2.4	2.7	0.3	0.2	0.2	9.0	9.0	0.4	0.1	0.4	1.0 2	2.0 0	0.7.0	0.2 0	0.3 0.2	2 0.1	67.2617	1
207741_x_at	NM_003293.2	tryptase alpha	175.0	99.7	2.2	2.2	1.8	0.5	0.5	0.3	0.2	9.0	0.2	0.3	0.3 2	2.2 0	0.7.0	0.1	0.5 0.3	3 0.6	59.015	15
207134_x_at	NM_024164.2	tryptase beta	214.6	112.3	3.1	3.8	1.7	1.2	0.3	8.0	9.0	0.3	9.0	0.4	0.1 0	0.5 0	0.2 0	0.1	.1 0.	4 0.2		æ
205653_at	NM_001911.1	cathepsin G	91.2	97.0	2.7	1.1	6.0	6.0	0.4	3.1	8.0	0.3		1.6	0.6 0	0.7.0		0.4	.3 0.3		_	49
205266_at	NM_002309.2	feukemia inhibitory factor complement protein CB namma	2.9	80 G	0.2	0.5	0.3	0.3	0.0	0.2	0.2	0.1	0.2	0.4	0.7 0	0.1	0.2 0	0.0	0.0	0.0	33,806	629
211743_s_at	BC005929.1	major basic protein	74.3	7.07	1.0	4.4	2.7	0.1	0.7	9.0	9.0			0.7				0.2 0	0.3 0.	2 0.3	31.5959	69
211549_s_at	U63296.1	15-nydroxyprosiagiandin dehydrogenase	48.3	48.3	2.1	2.5	1.4	1.5	£.	9.0	6.0	0.1	0.1	0.2	0.1	1.1	0.5 0	0.8	0.5 0.5	5 0.0	24.8454	22
206726_at	NM_014485.1	prostaglandin D2 synthase	119.0	94.0	2.0	8.7	5.7	0.7	0.1	1.0	0.5	0.2	0.2	0.2	0.6	3 0	0.7.0	.3	.2 0.	3 0.0	17.432	35
205011_at	NM_014622.1	loss of neterozygosity. 11. chromosomal region 2, gene A	76.1	6.07	3.0	6.8	4.5	2.3	1.0	1.4	1.9	0.2	1.1	1.2	1.8 2	2.9 2		نة _	.7 2.0	0 2.6	16.2511	=
205428_s_at	NM_001740.2	calbindin 2	16.7	63.4	1.3	7.0	6.0	0.4	1.6	0.7	1.3	1.8	1.6	3.6	1.5 1	7	0.7 1	1.1	.3 0.	6.0 9	_	-
219225_at	NM_024554.1	FLJ11413 typeine kinase with immunoolohulin	10.0	9.5	0.1	0.1	0.1	0.2	0.1	9.0	9.0	0.1	9.0	0.1	0.2 0	0.1 0	0 9.	0 1.0	0.0 9.	0 0.2	15.639	39
		and epidermal growth factor homology																				
204468_s_at	NM_005424.1	domains	3.6	2.8	0.5	0.3	0.1	0.0	0.1	0.2	0.0	0.0	0.0	0.1			0.0	0.0	1.1 0.	0.0		29
208343_s_at	AF 146343.1	CYP7A promoter binding factor	1.5	6.0	0.0	0.0	0.3	0.1	0.0	0.1	0.0	0.1		0.2	_		_	0	_			22
205051_s_at	NM_000222.1 R	CD117 c-KIT	92.1	82.8	6.2	8 .	7.3	2.3	5.2	6.5	2.7	0.7	2.1	8.0	1.1	0.8	0.2 0	0.5	0.1 0.1	1 0.5	12.3815	15
210102_al	BC001234.1	loss of neterozygosity, 11, chromosomal region 2, gene A	40.9	41.1	8.8	3.7	4.5	0.3	9.0	0.3	9.0	0.7		0.2	0.6	8.1	.8	1.7	.2			19
210796_x_at	D86359.1	sialic acid binding Ig-like lectin, siglec6	17.8	26.0	1.5	1.0	1.1	1.2	1.5	1.3	1.4	1.4	2.3	1.7	1.3 2	2.0	4.	1.5	1.8 1.0	9.0	10.5398	98
206519_x_at	D86358.1 R	sialic acid binding Ig-like lectin, siglec6	3.4	7.9	0.0	0.3	0.4	0.0	0.0	0.0	0.1	0.0	0.1	0.5		0.5 0	0.0	0.0	0.0 0.4			5
206480_at	NM_000897.1	feukotriene C4 synthase	8.8	16.0	0.3	0.5	9.0	2.4	9.	0.5	0.1	0.5		0.1								.
206617_s_at	NM_002910.4	renin-binding protein	10.6	6.2	1.2	. 0.1	0.4	2.8	1.6	0.5	0.1	9.0		0.1					9			బ
208089_s_at	NM_030794.1	tudor domain containing 3	0.0	13.4	0.8	9.0	0.5	0.5	0.7	0.5	0.5	0.1		0.3	0.0		0.7	-	9.0		_	97
205466_s_at	NM_005114.1	heparan sulfate 3-O-sulfotransferase ADAMTS3 a disintegrin-like and	- 1 8.	7.2	0.1	0.4	0.1	0.7	0.1	0.3	0.5	0.4	0.1	9.0	0.4	0.1	0 0.	0.0	4.1	0.0	8.15381	<u>.</u>
214913_at	AB002364.1	metalloprotease (reprolysin type) with thrombospondin type 1 motif, 3	6.8	4.3	0.5	0.7	8.0	0.5	7.0	9.0	9.0	0.5	0.4	0.7	1.2 0	0.7	9.6	.5	0.	5 0.5	7.2214	4
201860_s_at	NM_000930.1	tissue-plasminogen activator	22.2	29.7	0.5	0.4	0.1	0.1	0.3	0.5	0.5	0.1	0.1	1.3	0.4 1	9	0.1).2 (.6 0.	1 3.6	7.10135	35
206520_x_at	NM_001245.1	sialic acid binding Ig-like lectin. siglec6	19.0	22.7	2.4	1.1	9.0	1.1	1.5	1,3	1.0	2.0	2.0	0.4	1.4 2	2.9	2.1 2	2.3	1.9 1.9	9 1.2	-	33
220532_s_at	NM_014020.1	LR8 protein	52.4	18.2	8.0	9.0	1.0	6.4	5.9	1.5	1.2	0.4	0.3	1.4	0.6 2	2.2 (0.6	0.2	5.0 0.1	1 0.0		23
218169_at	NM_018052.1	FLJ10305	6.2	14.6	0.3	0.2	9.0	0.4	0.3	0.3	9.0	9.0	0.5	0.1	0.1 0	0.5	0.2	4	.6 0.	3 1.0	6.04597	97
221728_x_at	AK025198.1	nuclear receptor subtamily 1, group I, member 3	5.6	5.5	9.0	6.0	1.0	0.4	3.6	6.0	9.0	0.7	9.0	4.4	0.9 0.	6	0.8	0.7 0	0.7 0.	8 0.5		88
214028_x_at	AU156998	tudor domain containing 3	3.2	1.9	6.0	0.0	0.4	0.8	0.3	0.4	0.7	9.0	9.0	0.1	0.1 0	0.8	1.2	·	0.1 0.7	0.0	5.10738	88
221552_at	BC001698.1	fipase	10.6	3.4	6.0	1.8	1.1	6.0	1.1	1.0	6.0	0.2	0.2	0.7	0.1 0	0.1	0.2 0	0.7	0.9 0.3	3 1.1	5.02434	8
203367_at	NM_007026.1	MKP-1 like protein tyrosine phosphatase (MKP-L)	15.0	45.4	1.2	6.0	0.8	6.0	6.0	8.0	0.1	0.2		0.4	2.0 0		1.0		0.5 1.4			14
206997_s_at	NM_004807.1	heparan sulfate 6-0-sulfotransferase	4.2	3.8	0.2	0.1	0.1	0.4	0.4	0.1	0.3	0.3	0.4	0.3	ŀ	0.3	1	9.0	0.3 0.	65	4.81127	اء

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Prope ser	Accession #		Leanscripts	0000	E I		Strail	Small		702	Silidili	Surgill Surgill	-	١	ı	Sugn	ı	ı	ı	ı	١	h	
211517_s_at	M96651.1		IL-5R interleukin 5 receptor alpha	9.0	0.1	1.8	28.2	17.9	16.4	25.3	32.6	29.2	0.0	0.7	0.0	1.0	0.0	6.4	0.5	0.0	0.2	0.0	61.956
210744_s_at	M75914.1	œ	IL-5R interleukin 5 receptor alpha	0.2	1.6	15.4	30.8	15.3	18.7	40.2	29.7	33.0	0.3	0.2	1.5	0.5	0.1	.0.1	0.1	0.1	0.2	0.1	42.811
			CKIRC Chemoattraciant																				
206361_at	NM_004778.1	GPR	expressed on Th2 cells	4.8	0.3	22.0	40.5	15.3	33.7	37.9	38.9	42.3	1.0	. 2.6	2.2	2.1	8.0	1.4	1.0	1.2	6.0	0.5	16.642
206207_at	NM_001828.3		Charcot-Leyden crystal protein	1.1	0.1	270.0	219.1	203.6	226.8	233.8	179.7	163.1	2.1	19.4	19.1	49.3	2.8	1.0	4.4	6.0	2.0	0.4	15.164
203638_s_at	NM_022969.1	α	fibroblast growth factor R 2	0.5	0.1	7.3	43.5	33.0	4.	8.8	12.3	23.3	0.1	0.1	0.3	0.3	0.1	0.1	0.2	0.1	0.1	1.0	13.357
			egt-like module containing,	·																		_	
207111 at	NM 001974.1	GPR	sequence 1 (EMR-1)	1.5	0.1	16.4	49.5	34.5	85.9	93.1	91.2	93.5	2.7	3.7	5.4	5.1	3.4	1.7	9.0	7.1	1.6	0.5	8.0092
215248 at	AU145003		FLJ11581 fis, clone HEMBA1003598	0.7	. 0.3	5.7	4.7	5.2	4.5	2.1	2.8	2.1	1.0	0.1	0.1	0.1	0.2	0.1	0.1	0.1	0.1	0.5	7.2618
ı			dachshund (Drosophila) homolog								,												;
205471_s_at	AW772082		Hs.63931	0.1	9.0	-	7.	3.7	6 .4	œ ??	0.6	9.7	0.8	6.0	.	8.0	0.1	0.1	0.3	0.5	0.0	0.1	6.8198
218857_s_at	NM_025080.1		FLJ22316	1.0	7.0	18.0	35.3	18.6	13.7	12.5	16.3	15.4	1.3	1.7	6.0	1.9	0.3	0.5	0.4	2.4	0.5	0.5	6.5045
221169_s_at	NM_021624.1		histamine receptor H4	0.5	0.5	4.5	12.5	8.9	1.9	5.6	4.2	2.9	0.5	9.0	Ξ.	9.0	0.4	0.7	0.5	0.7	0.0	0.3	5.9783
201769_at	NM_014666.1		enthoprotin	6 9.	-6	41.7	101.5	. 0.9.	. 56.8	34.1	40.5	35.7	3.0	5.9	3.6	4.2	4.5	6.3	6.9	8.0	9.0	6.7	5.1015
			secreted throblast growth factor	,	,		,	. 6			,			,	;		;		,	,	4	-	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
208228_s_at	M87771.1		receptor (K-sam-III)	0.3	4.	3.7	10.2		3.9	6.9	4 .3	 	9.0	4.4		5.5	<u>.</u>	Ç.) (C	D.3	C	-	4.9922
213605_s_at	AL049987.1		hypothetical protein, MNCb-4779	8.0	0.0	10.8	46.8	44.7	6.7	. .	25.8	9.02	د .	4.4	4.5	4.2	5.9	7.	2.1	3.7	2.0	0.3	4.7426
205382_s_at	NM_001928.1		adipsin	3.5	0.1	62.0	206.0	176.5	62.4	48.1	60.5	44.8	7.1	32.5	20.5	16.3	0.1	9.0	0.1	15.6	0.1	3.8	4.7202
49452_at	A1057637		hypothetical protein LOC283445	9.0	0.2	5.2	7.7	9.9	6.9	5.9	10.2	12.7	1.1	0.2	0.3	1.3	0.1	1.6	1.6	0.5	1.3	0.8	4.7183
- 100000	1 000000	c	CD244 natural killer cell receptor	4	Š	,	04.0	7 33	c	9	3 96	0	ć	Š	;	,	,	Š	3.5	6.3			4 6207
22030/_at	NM_U16382.1	r	ZB4	C.3	3	2.0	9.70	4.00	ņ	9	0.02	0.0	5.	Š	Y.	3.0	Š	• 5	5.5	7.	-	- -	4.023
210108_at	BE550599	S	L type, alpha 1D subunit	0.2	0.1	1.0	1.7	2.4	0.0	0.8	2.4	4.8	0.0	9.0	0.5	0.5	0.0	0.3	0.1	0.0	0.3	0.0	4.5958
1 00000			protein kinase-related oncogene	9	;	613	,,,,	,	,	•	9	40	•	0	,	18.4	•	12.1	120	9 7			4 1083
209193_at	M24 / / 9.1		(PIM1)	Ð.	:	5.15	6.011		9.74	è				0.0		<u>.</u>	-	<u> </u>	<u>.</u>	D.	?	<u>.</u>	1.1303
			polyphosphate-1-phosphatase																			_	
202794_at	NM_002194.2		(INPP1)	5.7	6.2	7.1	25.8	20.5	17.9	24.8	53.6	57.8	2.8	2.7	5.6	4.0	1.9	4.8	5.6	4.8	1.9	8.8	4.1158
700000	MM 004007 4	Ċ	CCR3 chemokine (C-C motif)	ć	;	7	473	463.3	. 4	4	4	107	, 00	6	26.3	1 30	ç	2	Š	,	,		2 0525
208304_at	NM_001837.1	5	receptor 3	7.0	: ;	3. 10	146.9	102.2	40.4			6. 5	6.03	0.0	6.03	- 67	, .	5 1	• ·	7.	y !	; ;	0.0000
206111_at	NM_002934.1		eosinophil-derived neurotoxin	2.2	2.0	33.7	71.4	38.4	104.7	80.8	47.7	45.1	2.0	2.3	 -	89	7 .	0.7	0.4	14.8	0.7	0.5	3.7641
43427_at	AI970898		hypothetical protein LOC283445	0.8	0.7	2.1	4 .	4.3	 -	4.6	8.0	6.4	1.0	0.8	0.7	0.8	7.	1.0	<u></u>	9.0	Ξ:	<u></u>	3,7554
			inositol							;	:			1	į			•	٠.	:	,		0000
213804_at	A1039084		polyphosphate-5-phosphatase, 75kD	9.0	1.2	4.3	8.0	8 9.	9.6	2.6	6.3	9.7	0.7	0.5	2.2	1,7	4.	1.2	7.	1.0	E:	<u>.</u> ن	3.6088
. 209906_at	U62027.1	GPR	C3a receptor	12.4	11.2	44.1	72.3	20.2	18.8	38.8	70.3	29.9	1.2	1.2	1.8	3.8	9.1	1.6	1.5	3.0	9.0	0.5	3.6039
	07200114		ATP-binding cassette, sub-family C	0	•	,	9	9	;	6		700	~	36	,	4	9	4	7				3 5673
202804_at	AISSE/ IO		(CFIRMRP), member	0.0		7.7	7.0		7.5	70.0			3 6	2 1	;	3	9 9	;	,	; ;	? ;	; ;	9 6
221675_s_at	AF 195624.1		cholinephosphotransferase 1 beta	0.7	9	31.6	. ·	22.8	6.07	32.6	5.0	5.5	7.6	9	e i	D .	6.4	<u>.</u>	o	· .	9 9	, ,	3.3020
201562_s_at	NM_003104.1		sorbitol dehydrogenase (SORD)	2.0	1.5	5.9	4.2	4.	7.4	1.6	œ •••	1.2	1.7	5.2	1.5	1.3	1 .3	4.	9.	1.2	1.8	-	3.5098
210230_at	BC003629.1		FLJ23438 fis. done HRC13275	1.1	0.1	8.3	1.0	13.3	2.4	5.1	5.	4.1	0.1	0.5	0.5	0.7	0.3	0.5	9.0	0.8	.9	0.5	3.2532
219919_s_at	NM_018276.1		FLJ10928	0.1	0.5	3.1	1.8	2.5	1.7	4.3	1.7	2.1	0.1	0.5	9.4	0.3	0.1	0.1	0.1	0.1	0.1	0.7	3.2497
204301 at	NM 014867.1		KIAA0711	9.0	0.1	3.8	14.3	10.7	5.3	8.9	12.6	10.5	1.2	9.0	1.6	1.6	1.0	8.0	1.2	5.6	8.0	0.1	3.2398
			Grb 10- and Grb-IR-related splice																				
210999_s_at	U66065.1		variant 1	3.6	1.7	8.4	13.6	1	7.3	4.6	8.	6.4	2.2	1.7	6.	5.0	1.9	7	7.	1 .	0.4	9.	3.1044
200530 at	D25304 1		RacCdc42 guanine exchange factor	13.1	24.8	28.8	7.4.7	73.0	8	35.5	92.9	9 65	6.5		18.5	13.7	3.1	10.8	12.9	7.1	8.7	-0.	3.0943
- CCCCC2	1,0001.			;	;							7			47.6	4	36	:	42.7	*		47.0	2000
Z08921_s_af	L12387.1		Sorcin (SKI)	7.61	\$.	7.07	88.0	0.70	9.78	47.4	60.3	60.4	G	n O	9.	6.1.3	۷.۵	=	,	3		•	3.0020
			sulfurdaseadenosine																				
209043_at	AF033026.1		5-phosphosulfate kinase	19.9	16.1	39.0	87.1	75.2	66.4	68.7	89.4	71.4	13.8	20.0	46.5	8.9	6.3	6.7	4.4	10.6	11.0	22.7	3.0385

F. Eosinophil and neutrophil-selective transcripts (1/1).

Probe set	Accession #		Transcripts	MC. cord blood	Ind.	8 -	Ba 2 (small)	Ba 3 (small)	E0 1	Eo 2	Eo 3 (small)	Eo 4 (small)	Š +	2 Ne	Ne 3 (small)	Ne 4 (small)	=	CD4	8Q3	CD14	CD 19	æ	Eo+Ne S.L
221345_at	-	GPR	GPR43, PAR1-like	0.1	9.0	6.0	9.0	9.0	7.0	16.6	15.4	10.5	49.7	45.7	22.5	23.0	0.8	0.1	0.1	0.7	0.4	0.1	21.742
212860_at	BG168720		zinc finger, DHHC domain containing 18	2.2	0.8	4.1	3.9	4.0	16.0	17.4	14.8	16.7	59.2	53.9	30.9	39.8	1.1	2.4	3.8	5.6	1.5	1.3	6.5842
211576_s_at	BC003068.1		solute carrier family 19 member 1	1.4	0.1	0.7	1.4	0.7	5.1	5.9	1.5	12.4	19.8	18.7	27.1	28.1	1.8	6.0	9.0	2.7	0.5	0.8	4.99
214321_at	BF440025		nephroblastoma overexpressed gene ARF-GAP, RHO-GAP, ankyrin repeat	1.3	1.0	0.8	0.2	0.7	6.2	7.2	17.0	13.7	4.5	7.6	10.7	11.3	0.1	0.1	0.0	0.1	0.1	1.8	4.9823
218950_at	NM_022481.1		and plekstrin homology domains-containing protein 3	1.4	3.1	2.4	3.7	2.1	15.5	15.9	18.3	15.5	28.0	24.3	15.9	20.6	0.7	0.2	8.0	3.9	6:0	1.0	4.829
205681_at	NM_004049.1		BCL2-related protein A1	0.5	4.9	1.0	3.2	2.4	52.4	40.2	35.9	30.5	46.5	49.2	35.5	33.5	1.5	2.2	6.1	8.3	4.3	0.5	4.7782
203765_at	NM_012198.1		grancalcin	5.0	1.8	4.5	11.7	8.3	27.5	39.8	46.7	34.8	71.6	85.2	80.5	77.9	1.0	1.0	1.2	12.1	3.1	0.5	4.4403
213241_al	AF035307.1		cDNA FLJ36416 fis. done THYMU2011053	5.0	2.4	9.1	1.0	1.0	33.2	37.1	68.7	40.6	63.2	77.8	89.2	80.9	3.0	3.0	8.8	13.5	4.1	7.8	4.2603
221815_at	BE671816		hypothetical protein PRO2831	1.2	1.4	1.3	0.5	1.6	10.1	8.8	11.2	6.4	5.0	6.2	4.0	3.8	1.6	9.0	0.2	1.4	0.9	9.0	4.1325
			homolog of yeast tong chain polyunsaturated fatty acid elongation																				
214153_at	BE467941		enzyme 2	1.0	9.0	1.0	2.0	3.1	9.1	10.4	18.2	16.6	9.7	9.7	14.5	15.2	9.0	6.0	1.2	0.7	2.3	0.1	4.0797
212821_at	AU147160		KIAA0599	0.1	0.1	0.4	1.2	0.9	4.6	5.8	4.7	3.4	4.4	3.5	0.0	7.9	0.4	0.5	0.4	0.3	0.0	0.5	3.727
			Edg4, endothelial differentiation, fysophosphatidic acid																				
206723_s_at	AF011466.1 G	GPR	G-protein-coupled receptor, 4	1.2	1.4	2.1	3.6	5.6	13.7	8.0	19.6	22.3	24.6	17.0	26.1	29.1	0.1	3.6	2.0	3.8	1.9	0.9	3.7119
212360_at	AI916249		adenosine monophosphate deaminase 2 (isoform L)	1.0	2.0	5.6	5.9	4.6	26.6	18.7	30.2	31.9	78.7	63.1	79.5	91.3	3.4	5.8	6.1	12.6	3.7	5.8	3.5888
1 000000	10000111		transforming, acidic coiled-coil	Š	3	•	ć	6	ć	;		,	1	•		ć	?	;	4	9	;	,	0 4760
218308_at	NM_U00542.		containing protein 3 (TACCS)	Š	Š	0	0.0	9	3	y.	9	.	0	6.0	5.5	6.23	7.4	7	<u>.</u>	9.	2	3	5.4103
212629_s_at	AK023692.1		protein kinase C-like 2	1.9	9.0	 -	3.7	5.	-	9.2	17.7	13.8	16.7	23.6	27.0	78.4	7.	7.	2.2	4.2	4.2	5.9	3.2963
201739_at	NM_005627.1		serumglucocorticoid regulated kinase (SGK)	30.2	25.2	1.9	18.5	25.4	114.1	116.0	150.9	164.8	60.2	17.1	105.8	156.3	1.1	3.7	0.7	34.6	1.6	18.9	3.2456
209473_at	AV717590		ectonucleoside triphosphate diphosphohydrolase 1	1.4	9.0	2.1	1.6	2.0	15.9	22.3	39.5	29.1	14.1	13.7	23.7	13.8	5.6	5.6	1.3	6.5	8.8	9.0	3.0752
209304 × at	AF087853.1		growth arrest and DNA damage inducible protein beta (GADD45B)	2.0	0.3	2.7	4.	3.6	15.5	11.9	16.3	34.4	7.0	12.0	13.5	9.4	4.0	ω, *-	4.2	4	3.7	9.0	3.0295
210666_at	AF050145.1		iduronate 2-sulfatase	0.1	0.3	1.3	0.5	1.7	7.1	2.9	3.3	3.8	6.1	5.1	5.2	10.8	0.4	0.1	6.0	0.7	0.4	0.1	3.0005

G. Basophil and neutrophil-selective transcripts (1/1).

Ba+Ne S.I	5 8139R		4.97025	4.61523	4.57355	4.33664	4.14372	000433	3.891//	3.8469	3.84207	3.7116	3 48188	00.00	3.4769	:	3.401	3.31991	3.29317	3.18834	3.1591
	٦	_	5.5	-	0.5	8.0	£.	_;	- - -	4.5	0.3	1.2	8	?	0.7		5.5	6.0	0.7	8.0	0.1
CD 19	2 2		9.0	2.5	9.0	0.5	<u>2</u> .	,	0.0	5.6	1.0	Ξ.	,	5	1.0		2.3	1.7	0.4	0.1	5
		ì	0.0	3.0	5.6	2.3	9.0	,	9.0	7.3	4.3	2.3	4	?	3.6		4.6	2.4	3.3	0.1	5:
יט	1		9.	Rύ	1.1	0.2	1.3	;	 	5.	6.0	5.	a	9	9.		<u>.</u>	8 9.	9.0	0.1	1.5
70	1		0.5	2.0).5 ().1	0.8		6.4	2.9	0.8	4.			2.3			.3	0.2 (0.1	1.0
- -	۱,		7.	7.0	1.6),1	0.5		0.7	5.3	0.4	1.5			2.4			. 2.0	0.7	0.5	. 6.0
=	1		27.7	22.5	37.0	0.6	7.6		2.5	22.3		15.4	9		22.8		40.6	12.3		56.1	3.2
]_		_			11.2	8.4					13.3			23.5				5.3		3.3
Ne 3	Т			24.5	61.7	Ξ				25.7				-						60.7	
Ž c	7,8		14.0	24.4	32.8	6.7	10.2			23.0	14.5	11.7			14.8		30.7	6.5	16.7	14.4	8.2
Š ÷	දි		13.7	15.5	31.4	4.5	8.4	•	15.8	13.6	19.6	8.8	34.3		17.6		29.5	9.8	19.7	18.8	7.6
Eo 4	2,5	ì	4.2	3.8	9.0	1.2	9.	•	4	1.1	0.1	3.3	;	?	4.7		9.5	2.7	1.4	14.5	1.5
Eo 3	3.3	;	4.5	4.3	1.3	5.9	1.4	•	4.0	13.9	1.6	5.7	;		3.3		10.0	3.9	4.1	16.1	1.5
С	,	?	3.7	5.5	9.0	1.7	1.7	;		7.7	0.3	1.9	9	0.0	2.4		9.9	5.	3.0	5.9	6.
	36	ì	2.4	3.6	0.5	2.4	8.	,	1.2	4.0	0.3	2.1	,	3	3.3		4.9	5.0	3.2	2.1	1.5
Ba 3	47.3	?	18.8	19.2	25.4	26.4	4.7	;	6.3	70.8	20.9	13.2	;	32.4	9.5		22.9	8.0	8.7	34.3	5.8
Ba 2	16.3	?	23.5	16.2	13.7	26.2	5.5	;	10.4	82.2	18.9	16.2		6.10	8.3		22.8	7.6	11.0	27.9	4.3
Ba -	- :	!	10.1	4.2	8.5	5.0	4.5	,	3.9	28.8	5.5	4.3	9	0.0	4.8		10.4	3.9	13.8	4.4	5.0
M C	2		0.3	1.7	1.4	0.1	6.0		0.1	3.0	0.1	1.7		o.	0.4		2.4	9.0	0.1	0.2	9.0
MC, cord	1	?	1.0	5.4	0.1	9.0	1.0		0.1	5.0	0.1	2.5	;	4.7	0.4		2.2	0.4	0.3	0.9	0.0
Texas acids	COL 60 amonio		FLJ13386	kelch (Drosophila)-like 2	histidine ammonia-lyase	done 23551 mRNA	FLJ13676 fis	tumor necrosis factor (ligand)	superfamily, member 14 (TNFSF14)	mitogen-activated protein kinase 14	Hs.276590 ESTs	E2F transcription factor 3	autosomal highly conserved protein	(AHCP) transforming, acidic coiled-coil	containing protein 3	bromodomain adjacent to zinc finger	domain, 28	hypothetical protein PRO2198	clone 23551 mRNA	ring finger protein 24	FLJ13712 fis
3	Mar 046006 4	, 00000.	NM_025180.1	NM_007246.1	NM_002108.2	AF007132.1	AK023738.1		NM_003807.1	NM_001315.1	N54942	NM_001949.2	, 110010	NM_U16255.1	NM_006342.1		NM_013450.1	NM_018621.1	A1692428	NM_007219.2	AK023774.1
3	210730 et	210133_00	219242_at	219157_al	206643_at	213935_at	222151_s_at		207907_at	202530_at	217521_at	203693_s_at		203420_al	218308_at		203080_s_at	219999_at	213805_at	204669 s at	215555_at

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Transcripts	≥ & Ā	MC, cord N	MC. lung Ba 1	Ba 2 (small)	Ba 3 (small)	E0 1	Eo 2	Eo 3 (small)	Eo 4 (small)	å –	2 S	Ne 3 (small)	Ne 4 (small)	2	CD4 CD8	8 CD14	14 CD19	19 Fb	MC+Ba S.t.
carboxypeptidase A3	۲	137.1	1.0 107.6	6 139.0	173.1	2.5	1.6	2.8	1.9	0.1	1.4	2.0	12.4	0.2	1.2 0.	0.2 0).2	.5 0	0.5 59.198
TRK neurotrophin receptor		8.0	1.2 4.	6 9.3	8.1	0.2	0.5	0.0	0.1	0.2	0.0	0.1	0.1	0.1	0.1	-	.,	0.0	.0 34.7309
regulator of G protein signaling (RGS13)		6.2	8.4 5.	5.6 7.1	10.9	0.3	0.1	0.4	0.5	9.0	0.0	0.1	0.3	0.1	0.0	0.0	0.2	3.0	0.0 21.7762
		19.9	24.7 35.4	4 51.9	45.6	0.9	0.1	0.5	0.3	0.1	0.1	0.5	1.8	1.6	0.5 0.	2 0	0.0	3.3	.1 21.1783
solute carrier family 18		24.3	27.1 8.1	1 22.3	25.0	1.3	1.0	9.8	6.0	1.3	6.0	0.5	0.7	6.0	0.4	3).2	3.3	0.3 20.330
GATA-binding protein 2		18.4	12.8 55.0	0 31.7	24.7	2.0	1.6	1.9	1.7	6.0	1.9	1.3	3.7	5.6	0.8 0.	0.8	0.9	2 4.	.7 9.46893
Ec epsilon R I beta 15-hydmxxmstaalandin	_	0.7	28.6 34.9	9 12.6	8.9	1.5	1.0	0.8	0.8	1.9	6.0	1.7	1.5	1.7	0.7 0.	0.7 0).1	0.3	0.5 9.39967
sding frame		63.9	51.4 24.8	8 36.0	23.3	.	4.6	4.6	2.6	1.3	6.0	£.	1.2	6.0	₹.	5:1	7.4	0.8	0.4 6.14285
,		7.4	4.2 4.	7.4	9.9	1.2	1.6	2.0	2.4	6.0	1.9	1.1	2.1	1.1	1,4 1,	4	ri.	7.4	.6 3.3802
protein kinase, X-linked		6.2	5.2 5.2	2 12.8	16.9	2.3	2.7	2.5	5.4	9.0	0.4	0.7	1.2	1.9	1.5 2.			2.0	3.2606
low density lipoprotein receptor	-	24.6	16.8 19.		•		1.0	9.	6.0	8.0	5.4	-5	5.5			3.4	3.5		.3 3.1679

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Specific transcripts markers for non-granulocytes	MC	cord	
I. Specific transcripts markers for non-granulocytes	WC	cord	
I. Specific transcripts markers for non-granulocytes	MC	cord	
 Specific transcripts markers for non-granulocytes 	WC	cord	

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			cord	ΨĊ.		Ba 2	Ba 3			Eo 3	Eo 4		z	89.	16.4						
	Accession #	Transcripts	poold	lung	Ba 1	(small)	(small)	Eo 1	Eo 2	(small)	(small)	Ne 1	Ne 2 (s	small) (s	small) F	7	204	CD8	2014	CD19	٩
203547_at	U47924 CD4	CD4	5.5	4.7	2.2	1.1	1.2	2.0	1.6	0.4	6.0	0.7	1.1	0.7	9.0	3.5	15.0	0.3	17.6	1.2	0.4
	AW006735	CD8	0.5	1.7	2.6	2.3	3.2	1.3	1.1	2.2	1.7	0.3	0.5	1.8	0.3	4.2	3.0	76.1	9.0	9.0	0.4
	NM_001770.1	CD19	0.7	0.1	0.7	. 0.4	0.7	0.8	0.5	0.5	0.2	9.0	1.5	1.1	0.7	4.0	0.0	0.1	0.1	19.1	0.4
	L14458.1	lgG V-J region	0.2	0.1	0.1	0.1	0.2	0.5	0.5	0.2	0.3	0.1	0.3	0.5	0.5	3.8	0.1	0.5	0.1	70.9	0.1
	NM_000591.1	CD14	18.4	2.4	0.3	0.1	9.0	8.8	9.0	1.1	3.2	40.7	49.6	58.7	68.2	5.9	9.4	0.0	97.2	1.1	6.0
	NM_005211.1	v-fms M-CSF receptor	3.8	0.5	9.0	0.7	0.3	3.0	1.1	1.7	1.5	8.8	8.7	6.3	6.4	5.2	2.8	0.7	42.1	1.4	6.0
	U63041.1	CD56	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.5	0.1	0.1	0.1	0.2	0.1	0.1	0.1	0.1	0.5	0.1	0.0
	M35999.1	CD61 glycoprotein Illa	1.7	9.4	0.5	0.1	0.3	0.1	0.1	0.1	0.1	0.1	0.4	0.1	0.2	48.3	0.1	0.3	0.1	0.5	0.5
	AK026737.1	fibronectin	0.3	1.5	0.3	0.3	0.5	0.1	1.0	0.3	9.0	0.1	0.1	0.2	0.1	0.3	0.1	1.0	0.3	0.1	97.6

J. Raw AD levels for the median values used to normalize the raw AD levels, and the housekeeping genes.

			MC,										,								
	Accession		cord	MC,		Ba 2	Ba 3			E0 3	Eo 4			Ne 3 N	Ne 4						
Probe set	*	Transcripts blood	poold	Buni	Ba 1		(small)	Eo 1	Eo 2 ((small)	Ne 1	Ne 2		(small)	_	8	800	CD14	CD 19	٩
AFFX-HSAC07/X00351_3_at	X00351	beta-actin	18534	19383	14638	20922	19151	22019	18638	21153	18568	19406	22302	16589	. 695/1	18295	18782	20805	20661	17542	20399
AFFX-HSAC07/X00351_M_at	X00351	beta-actin	22898	21777	12940	11533	1608	22560	18785	14861	13431	23106 2	23373	6861	8019	19368 2	20362	22761	22294	19087	22757
AFFX-HSAC07/X00351_5_at	X00351	beta-actin	15642	15838	9516	2927	2066	17186	21061	4861	3771	. 21181	19768	2165	2193	14887	16381	18218	17583	15063	16795
AFFX-HUMGAPDH/M33197_3_at	M33197	САРОН	14906	13632	3640	10477	9332	2649	2844	5498	2605	2549	3186	3586	3538	5929	6429	8609	9756	5180	19674
AFFX-HUMGAPDH/M33197_M_at	M33197	GAPDH	15891	16852	3218	7887	6963	2382	2565	3720	1838	2194	2796	2312	2154	6182	5931	5921	11464	4344	17013
AFFX-HUMGAPDH/M33197_5_at	M33197	САР DН	16298	16701	3479	4559	5110	1827	3121	2127	1168	2492	3476	1636	1355	4655	9889	6260	9905	5636	23350
The median value of 22283 transcripts			121	169.3	85.3	112.2	110.9	93.7	102.1	127.3	114	62.7	73.1	87.4	87.3	82.3	152.5	121.3	107.1	114.1	183.8

Abbreviations used in the table A-I were (small); the results obtained by the small sample protocol (see materials and methods), R; receptor, and ICN; ion channel.